

<u>ARCS II PROGRAM</u>

Remedial Planning Activities at Selected **Uncontrolled Hazardous Substance** Disposal Sites Within EPA Region II (NY, NJ, PR, VI)

> **FINAL** SCREENING SITE INSPECTION (SSI) CAPTAIN'S COVE CONDOMINIUM SITE GLEN COVE, NASSAU COUNTY NEW YORK

> > SEPTEMBER 1995

VOLUME V OF V

EPA Contract 68-W8-0110

An ENSERCH® Engineering and Construction Company

EPA WORK ASSIGNMENT NO: 076-2JZZ EPA CONTRACT NO: 68-W8-0110 EBASCO SERVICES INCORPORATED ARCS II PROGRAM

FINAL
SCREENING SITE INSPECTION (SSI)
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NOTICE

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RADIOACTIVE CONTAMINATION AT

CAPTAIN'S COVE CONDOMINIUM SITE

The radiologically contaminated soils at the Captain's Cove Condominium Site should be remediated in accordance with current efforts underway at the Li Tungsten Site, a National Priority List (NPL) site.

The following presents relevant points regarding the similarities between the two facilities.

Historically a portion of the Captains Cove Condominium Site was used as a municipal sanitary landfill by the town of Glen Cove beginning in 1971 (Ref. 2, p. 10 of 16). It was speculated that the site was also used for disposal prior to this documented date (Ref. 3, p. 28 of 39). In the same time frame (the 1940s through the early 1970s), the nearby Li Tungsten facility processed tungsten ores with elevated concentrations of uranium and thorium (Ref. 4, p. 6 of 23). Anecdotal information coupling the sites was provided by a former employee of Li Tungsten (Ref. 5, p. 1 of 2). During a telephone interview the former employee noted that he was told by "old timers" that they routinely dumped waste slag at the Captain's Cove Condominium Site (Ref. 5, p. 1 of 2). Further, he stated that employees would take boxes or crates of material by fork lift down the road and dump the material at the Captain's Cove Condominium Site (Ref. 5, p. 1 of 2). This is similar to waste disposal at Li Tungsten, e.g., wastes were dumped in the parking lot near the main building, across the street by the woods, back by the reduction building and hydrogen tank (Ref. 5, p. 2 of 2).

The sites are located on the same road less than 1,000 feet apart (Ref. 6, p. 6 of 48). The land between the sites is a boat yard on which no known industrial processes occurred. The road is the only land access to both sites as they both abut Glen Cove Creek.

As regards to prior investigations, the source of the radiological materials at the Li Tungsten Site resulted from the smelting and refining of tungsten metal from ore materials principally Schelite (CaWO₃) from Canada and China that also contained concentrations of uranium and thorium as accessory metals (Ref. 4, p. 6 of 23). The smelting process extracted the tungsten metal and concentrated the uranium and thorium in the waste slag (Ref. 4, p. 6 of 23). Tungsten concentrations and other heavy metals in the waste are assumed to result from the fact that the refining process was imperfect (Ref. 7, pp. 19 and 20 of 27).

The concentrations of uranium and thorium in the tungsten ore and some thorium processing were sufficient to require Li Tungsten (and its predecessor the Wah Chang Trading Company) to acquire a Radioactive Materials License from the NYS Department of Labor i.e., and a United States Atomic Energy Commission Source Materials License i.e., a license for ores with concentrations equal to or greater than 0.05% by weight of uranium and/or thorium (Ref. 4, p. 7 of 23; Ref. 8, p. 1 of 2; Ref 9, p. 1 of 2).

The similarity of the radionuclides and concentrations on both sites was established through a review of prior investigations (Ref. 4, p. 6 of 23, Ref. 10, p. 28 of 70). On the Li Tungsten Site, concentrations of uranium and thorium in the input ores average about 10 pCi/g for all thorium

and uranium chain nuclides, with measured concentrations in ore and intermediate processed material at 23 pCi/g ²³²Th and 27 pCi/g ²³⁸U(Ref. 4, p. 20 of 23),. The range of concentrations in waste material increased considerably up to and greater than 1,000 pCi/g (Ref. 4, p. 6 of 23). Similar radiological concentrations were found on the Captains Cove site (Ref. 10, p. 28 of 70). Trench samples taken by Fred C. Hart indicated concentrations in the 10 to 100's of pCi/g of uranium and thorium (Ref. 10, p. 28, 29, 30 of 70). It should be noted that the samples at the Captains Cove Condominium Site were taken in the fill material (from the surface to about the 10 feet depth) (Ref. 10, pp. 29, 30 of 70).

The last characteristic is the result of the recent soil investigation conducted in April 1995 (Ref. 11, pp. 1 to 28 of 28). A total of nine surface and three subsurface soil samples, including the background samples CC-SS11-01 and CC-SS11-02 (surface) and CC-SS11-03 (subsurface), were collected for analysis. The radiological and metal (tungsten) samples were taken from areas exhibiting elevated gamma exposure which were previously identified as containing radiological anomalies (Ref. 11, pp. 28 of 28). The background samples were taken from the Garvies Point Preserve (Ref. 11, pp. 28 of 28). The background sample location is off the site about 150 feet perpendicular to the north western edge of the site by the second gate (Ref. 11, pp. 28 of 28). No industrial processes or major disturbances e.g., fill deposition were known to be conducted at the background location (Ref. 11, p. 28 of 28). Five soils samples were collected from the Li Tungsten Site (Ref. 11, p. 28 of 28). The samples were analyzed for metals, including tungsten, and for radionuclides of the uranium and thorium decay series (Ref. 12, pp. 1 through 39, Ref. 13, pp. 1 through 45 of 45).

Background Sample Results

Ref.13, pp. 24, 26, 43, 45 of 45

The background samples data indicate that natural uranium and thorium concentrations in this area are close to the average crustal abundance, with no evidence of elevated concentrations (Ref.13, pp. 24, 26, 43, 45 of 45). The crustal abundance is about 0.6 pCi/g for the members of the uranium series and 1 pCi/g for the members of the thorium series (Ref. 4, p. 6 of 23). Results of the background samples for the uranium series are presented in Table 1.

Table 1				
	²³⁸ U (pCi/g	²³⁵ U (pCi/g)	²³⁴ U (pCi/g)	²³⁰ Th (pCi/g)
CC-SS11-02	0.953 ± 0.166	0.0388 ± 0.0306 J	0.847 ± 0.155	0.926 ± 0.133
CC-SS11-03	0.454 <u>+</u> 0.114	0.0275 ± 0.027 J	0.505 ± 0.122	0.467 ± 0.166

As indicated above, secular equilibrium is evident for ²³⁸U, ²³⁴U and ²³⁰Th (Ref. 14, p. 3 of 3). Also the ratio of ²³⁴U/²³⁵U is 21.8 and 18.4 is close to the naturally expected abundance ratio of 22 for these two radionuclides (Ref. 14, p. 2 of 3).

The thorium series shows similar results in the background samples as presented in Table 2.

	Table 2	
	²³² Th	²²⁸ Th
CC-S11-02	1.22 <u>+</u> 0.153 J	1.13 ± 0.165 J
CC-S11-03	$0.445 \pm 0.114 \mathrm{J}$	0.471 ± 0.162 J

Secular equilibrium is evident between these two thorium chain members in both samples (Ref. 13, p. 43, 45 of 45).

Correlated Radiological and Tungsten Contamination

All the soil samples collected from Li Tungsten with radiological concentration significantly above background also have high concentrations of tungsten.

Table 3

	²³² Th (pCi/g)	²³⁸ U (pCi/g)	W (mg/kg)	
LT-SS01-01	10.1 ± 3.55 J	52.2 <u>+</u> 9.30 J	3050 J	
LT-SS02-01	17.7 ± 4.65 J	2 6.2 <u>+</u> 7.44	16200 J	
LT-SS05-01	24.8 ± 6.49	165 ± 18.6	1160	
LT-SS05-01D	25.1 <u>+</u> 7.32	154 ± 18.3	1420 J	

(Ref. 12, pp. 26, 27, 30, 31 of 39, Ref. 13, pp. 11, 12, 16, 17, 30, 31, 35, 36 of 45)

The same pattern of elevated concentrations is also observed in the Captain's Cove Condominium soil samples. That is elevated concentrations of uranium, thorium and tungsten.

Table 4

	²³² Th (pCi/g)	²³⁸ U (pCi/g)	W (mg/kg)
CC-SS14-01	20.0 <u>+</u> 5.71	18.6 ± 3.37	3200 J
CC-SS15-01	16.2 <u>+</u> 4.40	18.4 <u>+</u> 4.46	3820 J

(Ref. 12, pp. 20, 21 of 39, Ref. 13, pp. 22, 23, 41, 42 of 45)

This correlation strongly suggests that the ore and processed material was derived from a single source.

Uranium Decay Series Disequilibrium

Examination of the individual isotopes of the ²³⁸U decay series presents additional evidence that the radiological constituents found on the two sites may have a common source (Table 5). In naturally occurring undisturbed radiological materials, all isotopes of the decay series are in a state of secular equilibrium, that is, they are all present at the same activity (Ref. 14, p. 3 of 3). This situation is illustrated by the background samples previously discussed (Table 1 and Table 2). If the decay members are subjected to chemical or physical separation, secular equilibrium may be disrupted, allowing the various isotopes to be present at significantly different activities. If, for example, uranium was preferentially extracted with the tungsten from the original ore, the isotopic activity of ²³⁸U and ²³⁴U would be less than the activity of ²³⁰Th.

The tungsten refining operations at Li Tungsten included physical and chemical processes which included vibrating screens, magnetic separators, electrostatic separators, acid leaching, floatation and fusion (Ref. 7, pp. 15, 19, 20 of 27). Each of these processes has the ability to alter the initial secular equilibrium in the input tungsten ore. This is evident in the Li Tungsten samples as presented in Table 5.

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	²³⁸ U (pCi/g)	²³⁴ U (pCi/g)	²³⁰ Th (pCi/g)	
LT-SS01-01	52.2 ± 9.30 J	46.3 ± 8.72 J	11.1 ± 3.70	
LT-SS02-01	26.2 ± 7.44	10.7 <u>+</u> 4.94	20.0 <u>+</u> 4.96	
LT-SS05-01	165 <u>+</u> 18.6	155 ± 17.8	344 <u>+</u> 24.7	
LT-SS05-01D	154 <u>+</u> 18.3	148 ± 17.8	303 <u>+</u> 26.0	

(Ref. 13, pp. 11, 12, 16, 17, 30, 31, 35, 36 of 45)

The disequilibrium in sample LT-SS02-01 is tenuous and could be a statistical anomaly as physical metal recovery processes would not separate ²³⁸U from ²³⁴U.

Similar states of disequilibrium are also observed in radiologically contaminated samples collected from the Captain's Cove Condominium Site as presented in Table 6.

Table 6

	²³⁸ U (pCi/g)	²³⁴ U (pCi/g)	²³⁰ Th (pCi/g)	
CC-SS14-01	18.6 ± 3.37	23.9 ± 3.91	45.2 <u>+</u> 7.47	
CC-SS15-01	18.4 ± 4.46	20.8 ± 4.64	39.4 <u>+</u> 6.90	
(Ref. 13, pp. 22, 23	3, 41, 42 of 45)			

Ratios of the radionuclides ²³⁸U/²³⁴U, ²³⁰Th/²³⁸U and ²³⁰Th/²³⁴U corroborate the disequalibria (Table 7). The most highly contaminated samples from Li Tungsten (LT-SS05-01 and LT-SS05-01D) have essentially the same ²³⁰Th/²³⁸U and ²³⁰Th/²³⁴U ratios that are found in the contaminated Captain's Cove Condominium samples (CC-SS14-01 and CC-SS15-01). These ratios are significantly higher than those measured in the background soil samples CC-SS11-02 and CC-SS11-03, which exhibit secular equilibrium.

Table 7				
	²³⁸ U/ ²³⁴ U	$\frac{230}{\text{Th}} / \frac{238}{U}$	$\frac{230}{\text{Th}} / \frac{234}{\text{U}}$	
LT-SS05-01	1.06 ± 0.171	2.09 ± 0.279	2.22 ± 0.301	
LT-SS05-01D	1.04 ± 0.176	1.97 ± 0.289	2.05 ± 0.302	
CC-SS14-01	0.778 ± 0.190	2.43 ± 0.596	1.89 ± 0.440	
CC-SS15-01	0.885 ± 0.292	2.14 ± 0.640	1.89 ± 0.537	
Background Samples				
	Table 8			
CC-SS11-02	1.13 ± 0.284	0.972 ± 0.219	1.03 ± 0.276	
CC-SS11-03	0.899 ± 0.313	1.02 <u>+</u> 0.444	0.92 ± 0.320	

Thus the separation processes employed by the Li Tungsten facility are believed responsible for a similar departure from secular equilibrium observed in soil data from both sites.

The evidence presented above illustrates similarities between the uranium and thorium isotopic concentration and tungsten concentration found on the Captain's Cove Condominium Site and that found on the Li Tungsten Site. This particular combination of correlated concentrations can be explained by deposition of raw ore or intermediate or fully processed wastes from the Li Tungsten refining operation. As no similar refining process was conducted on the property designated as Captain's Cove Condominium Site, the existing information and similarities between the radiological contamination at both sites strongly suggest that the tungsten and radiological contaminants found in the Captains Cove Condominium samples originated from the adjacent Li Tungsten Site. This link provides a logical explanation for an otherwise rare combination of correlated isotopes and concentrations and thereby establish the similarities of the radiological wastes at the Li Tungsten and Captain's Cove Condominium Sites.

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REFERENCE 1

Sites Which Are Difficult to Address

One commenter said that "unbounded or unmanageable sites, such as well fields" should not be included on the NPL in response, EPA believes that unless a remedial investigation and feasibility study has been completed at a site, it is not possible to specify whether a site presents a manageable problem. Furthermore, at many of thos sites where commonly applied remedial actions are infeasible, some response actions short of waste removal or source controls, e.g., providing alternative water supplies, may be appropriate. EPA believes that the technologies for response actions have been developing rapidly; a response which was infeasible in the past may become feasible in the near future. Finally, with the case specifically mentioned, wellfields, the Agency has generally found the need for CERCLA response particularly acute since this generally involves contamination of public water supplies. Hence, EPA has not attempted to exclude sites which are especially difficult to address through current response technologies.

Noncontiguous Facilities

Section 104(d)(4) of CERCLA authorizes the Federal government to treat two or more noncontiguous facilities as one for purposes of response, if such facilities are reasonably related on the basis of geography or their potential threat to public health, welfare, or the environment. As previously stated (48 FR 65058, September 8, 1983), for purposes of the NPL EPA has decided that in most cases such sites should be scored and listed individually because the HRS scores more accurately reflect the conditions at the sites if each is scored individually. In other cases, however, the nature of the operation that crested the sites and, possibly, the nature of the appropriate response may indicate that two geographically separate properties should be treated as one site for purposes of listing. EPA has done so for some sites previously listed separately on the NPL

Factors relevant to such a determination may include whether the two or more areas were operated as parts of a single unit. Another factor is whether contamination from the two or more sites is threatening the same part of an aquifer or surface water body. Finally, EPA will also consider the distance between the noncontiguous sites and whether the target population (i.e., within 3 miles) is essentially the same or substantially overlapping for

the sites.

One commenter. Governor Bond of Missouri, submitted the 33 known dioxin sites in that State as a single site on the NPL. Using characteristics from various sites, he assigned a single HRS score to the 38 sites. Governor Bond maintained that the dioxin was produced by a single result generator and that the sites had a common method of disposal. According to the Governor, by treating the sites individually RPA has complicated negotiations for health studies.

development of cost recovery suits, and the State's accounting procedures.

EFA carefully considered the Covernor's proposal and, taking into account the factors discussed above, decided that his reasons did not warrant consolidating the 33 sites into a single The sites are dispersed over a wide area of the State and affect different larget populations. The 33 sites sensially comprised different disposal operations rather than parts of the same facility. Many of the 33 sites would not individually acore high enough to be on the NPL and, thus, the overall score for the 23 sites would be misleading. EPA has also concluded that listing the 33 wites as a single site on the NPL is not a prerequisite for developing a consulidated response strategy for the Missouri dioxin altes. Many of these altes may qualify for Fund-financed removal actions. The Agency is currently evaluating ways of coordinating possible response strategies at these sites to alleviate the problems which Governor Bond has

Another commenter expressed the view that my grouping of noncontiguous sites would be inappropriate. EPA disagrees. In some instances the property boundaries or other factors commonly used to define a site may not be very useful or reasonable for determining if a problem involves one alte or several. One example is the Minker/Stout/Romaine Creek site in Missouri where dioxin contaminated **soils were used as** fill in several yards in a residential neighborhood. Even though the contaminated areas are not contiguous and the properties involved bave several different owners, the Agency determined that the site was really a single operation, that the same target populations might be affected. and that there is no logic to support treating the various areas as separate sites: Given the many factors involved in making such determinations and the differing importance that each factor may take on in various situations, the ency must weigh each situation individually to determine if

noncontiguous disposal areas single site or several.

Where EPA determines, based above considerations, that two in noncontiguous locations are more logically considered as a single at they will appear actions, it is prescribe them; EPA may decide to response efforts should be distinguished appearately on the NPT single response if it appears costs affective to do so.

Scoring of Air Releases

A comment was received concern how past air releases are scored. Language in the preamble to the ENCP caused a commenter on the By Sorrell. Louislana site to question whether past air releases may probe included in a site's HRS score if issue is discussed in detail in the "Support Document for the revised "National Priorities List—1984" for Bayou Sorrell site. However, the may points of this issue are presented in following discussion.

EPA believes that past air relea included in a site's HRS score. The stipulates that a site is to be score an air release if data "show levels of contaminants at or in the vicinity of facility that significantly exceed contact background levels, regardless of the frequency of the occurrence (47 FR) 21236). According to the HRS as established in the NCP revision therefore, the single evidence of an release such as that which occurr Bayou Sorrell, requires that the scored as having an observed re air. This approach to scoring has clarified by EPA's stated policy sites are to be scored on the basis conditions existing before any remeasures were performed. This p was clearly stated at the time of promulgation of the NCP revisions FR 31188), and EPA considers it to firmly established as part of the H addition, the Agency has attemp clarify further the reasons for this in subsequent statements (48 FR 49

Several considerations underlied policy. Actions by States to conduct enforce cleanup might be discourant of a site could report a cleanup of a site could report a could r

Another concern is that responsed to comminmel, incomplete cleanup action sites that might reduce the HRS

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Fremedy the problems

To For example, a site may problems in all three routes—
water, surface water, and air.

To the sir route is remedied. In a site in the site in the partial site in the same the partial site in the same that the site would not so the basis of the latest so the site would not existing prior to the remedy if route (48 FR 4084).

The consideration is that the HRS signed according to the site approximations of risk that

ird consideration is that the HRS is fined according to the Hole approximations of risk that the derived from certain basic life is a site as they existed prior Cleanup actions. Where the data is reflect conditions after some site actions, the assumptions upon the HRS was designed may no the HRS was designed may no the impropriate, and the score polyrepresent an approximation that is accurate or consistent points for other sites. All three of considerations are explained in in the presemble to the initial NPL (19664-5).

FR 70864-5). Mother consideration is that the level return provided by the HRS and the ting process, while sufficient to de å general approximation of risks maparison among sites, is not whily detailed to evaluate the cy of cleanup actions. The HRS rigned to take into account as lactors regarding the condition of and the risk they present as can be the country. It does not take into ni factors that the Agency believes Ed require sophisticated data or in developing the HRS, EPA ered evidence that a release background has occurred is easify determined. However. ory determined that evidence as er past cleanup actions are and to have chiminated the release polectial for future releases is much difficult to obtain and evaluate.

after the NPL listing probes has identified a limited number of sittle as potential problems. Having taken this approach in the HRS, EPA must apply it consistently to individual sites.

A commenter on the Bayou Sorrell.
Louisiana site cited preamble language which states that "sir releases must currently exist, must be measured, and must not be caused by disturbances from investigations" (47 FR 31289). EPA believes that the commenter took this language out of context. Read in context, it in no way contradicts the Agency's policies of scoring on the besis of a single observation and according on the basis of conditions existing before any cleanup actions.

The portion of the procemble (47 FR 31189) containing this language was written in response to comments arguin that the HRS should provide for accring for the potential of a release, rather th only scoring when an actual release is observed. The HRS does score for potential releases in the ground water and surface water routes if no actual release has been observed. For the air route, however. EPA believed that evidence of the potential for an air release could not be easily established and would be too tenuous a possibility to warrant taking it hito account. Therefore, in order to calculate any score at all for the air route, an actual release must be observed. By sixting that air releases must "conjuntly exist." EPA was attempting to sixplain that the release must have actually occurred. rather than being merely a potentiality. This interpretation is consistent with the actual HRS instructions, which require "date that show levels of a contaminant at or in the vicinity of the lackity that significantly exceed background levels. regardless of the frequency of occurrence" (47 FR \$1236).

Any other interpretation of this language would be illogical. If the word "current" were to be interpreted as meaning "today." then an observed release to air would have to be

issues that were not site specific. General comments on the NPL are addressed throughout this preamble Significant comments regarding specific sites are addressed in the "Support Document for the Revised National Priorities List-1984." A number of the site-specific comments addressed similar issues, and EPA's rationale for addressing those issues is presented in this section. Many of the lasses raised in comments are the same as those raised reviously and discussed in the previous final rulemaking on the NPL (48 FR 40658. September 8. 1983). The Agency's positions on these issues remains unchanged.

Waste Quantity

A number of commenters said that the waste quantity values assigned under the HRS were too high because EPA had included the non-hazardous constituents of the hazardous substances in calculating the quantity of waste located at the facility. Commenters raised similar issues when the first NPL was published (48 FR 40658, September 8, 1983), and EPA's response remains unchanged.

Consideration of Flow Gradients

in some instances commenters maintained that, based upon their conclusions regarding prospective movement of contaminants in ground waters, the values assigned by EPA to population served by ground water are too high. The commenters said that EPA should only count the population using those wells which they believed would be affected by the releases. As was the case with the waste quantity issue, this issue was addressed and resolved when the NPL was first promulgated [48 FR 40658). The rationale for the Agency's approach is futher discussed in the preamble to the NCP (47 FR 31190-91 July 16, 1982) and is equally applicable now. The HRS specifies that all the population using the aquifer of concern within 3 miles of the facility should be

Federal Register / Vol. 48: No. 175 / Thursday, September 8, 1983 / Rules and Regulations

40663

consider eligible for listing on the NPL ration of this factor in the HRS. the Agency decided not to cevidence of frequency and mation of a release, as explained Promulaction of the HRS (47 FR To do otherwise would render Process unnecessarily end time-consuming. which vert funds from cleanup and impede the progress of the THE PA recognizes that these Tabons are very relevant to the risks presented by a site dedies, if any, that should be Grd Factors of this type. are intended to be evaluated

This would not only entail considerable, expense but would also allow the assignment of an observed release to the air to be negated by a removal or remedial action. The Agency has consistently scored sites on the basis of conditions before removal or remedial actions, as explained in 48 FR 40664.

VIL Changes From the Proposed NPL

The Agency received a total of 128 comments on the proposed NPL apdate. Of these, 112 comments pertained to 80 of the proposed sites. The remainder of the comments addressed sites that were not proposed or generic or technical

included in the calculations of population served by ground water. The Agency's approach is based on the difficulty of predicting precisely the movements of ground water based on the limited amount of data consistently available at the time of HRS scoring. Furthermore, in establishing the rating scales, the Agency took into account the fact that most wells within 3 miles would not be affected. If EPA were to establish rating scales based only on the populations that have been or are certain to be affected, the scales would have assigned high values for much smaller populations than those specified

TOTAL P.04

consider eligible for listing on the NPL those RCRA facilities at which a filicant portion of the release present to come from "non-regulated mits" of the facility, that is, portions of the facility that ceased operation prior is lausery 26, 1963.

Releases of Mining Wastes

Some commenters presented the view CERCLA does not authorize EPA to repond to releases of mining wastes, and that sites involving mining wastes should not be included on the NPL. This view is based on the interpretation that dining wastes are not considered hezardous substances under CERCL CERCIA includes in its definition of iazardous substances materials that constitute hazardous wastes under the Resource Conservation and Recovery Act (RCRA). In the 1980 amendments to RCRA, the regulation of mining wastes oder Subtitle C of RCRA was temporarily suspended and that Suspension is presently in effect. For bat reason, the commenters believe that mining wastes should not be considered bazardous substances under CERCLA. EPA disagrees with the commenters interpretation. The Agency believes that mining wastes can be considered chazardous substances under CERCLA if It meets any of the other statutory ecriteria (e.g., if the material is also a hazardous air pollutant listed under section 112 of the Clean Air Act). More importantly, however, EPA's authority to respond to mining waste releases. and the Agency's ability to list mining waste sites on the NPL does not depend on whether mining wastes are hazardous substances. Section 104(a)(1) of CERCLA authorizes EPA to respond to releases of not only "hazardous Substances." but also "any pollutant or contaminant." "Pollutant or contaminant" is defined very broadly in section 104(a)(2) to include essentially any substance that may cause an adverse effect on human health. EPA is convinced that mining westes can satisfy these minimal criteria. that the Agency therefore has the authority to respond to releases of mining wastes. and that listing of mining weste sites on the NPL is appropriate. Commenters also presented the view bat it is unclear whether CERCLA was intended to address the type of waste problem characterized by low concentrations and large volumes. associated with mining waste. They argued that the approach taken under RCRA of preparing a study of mining wastes before determining whether Egulation of such wastes is appropriate. ould be adopted in the CERCLA Program as well. Commenters suggested

that as a policy matter, long term permanent remedial actions could be postponed and only removal actions taken at such sites when emergency conditions warrant.

As described above, however, the response authorities of CERCLA are very broad. As long as EPA has the authority to respond, and no other Federal statute provides authority comparable to CERCLA, the Agency has the obligation at least to evaluate the precise extent of the risk and the possible response actions at all sites that upon preliminary investigation appear to present a significant risk. EPA should also remain free at least to consider all types of response actions at all sites in order to determine which is the most appropriate and cost-effective. and should not limit itself to considering only removal actions at a particular class of facilities. Inclusion of the NPL is appropriate in order to begin the process of determining how to address such sites. Since inclusion on the NPL does not determine whether response actions will be taken or what response is appropriate. BPA is free to develop an approach for responding to mining waste sites that takes into account any unique features of such sites.

Comments also presented the view that the HRS is not an appropriate tool to estimate the risk to health and the environment presented by mining waste sites.

They pointed out that the HRS does not consider concentration levels at the point of impact, but rather the mere presence of the substance in the environment. As explained in Part VII below, however, the purpose of scoring for an observed release without taking level of concentration into account is simply to reflect the likelihood that the subject substances will migrate into the environment, which in the case of an observed release is 100 percent. Future releases, or even current releases for which concentration data do not exist. may raise the level of concentration to the point that it presents a greater risk than the release first observed. While releases from mining waste sites may be somewhat less likely than releases of man-made chemical substances to ever reach extremely high concentrations. harmful concentrations can occur from mining waste sites and the distinction is not sufficient to invalidate the HRS as an appropriate model for scoring mining waste sites.

Another comment was that the locations of mining waste sites are generally rural, so that the only sizable target population are far downstream. The comment alleged that these

populations are considered in the HRS scoring but in reality may never be affected. This assumption, however, is false. The HRS considers only those persons living within a three mile radius of the site as constituting the target population. If a mining waste site has a high score for this factor, it indicates that despite the fact that the locations of such sites typically are rural, this particular site has a significant number of people within three miles.

Indian Lands

EPA has always considered sites on Indian lands to be eligible for inclusion on the NPL. However, one commenter was concerned that some sites on Indian lands may not have been included in the State evaluation of NPL candidate sites because Indian lands are not subject to State jurisdiction. The Agency recognizes that this may happen. However, EPA Regional Offices may also evaluate sites for inclusion on the NPL. The Agency urges commenters to submit information on any sites which they feel may not have been evaluated during preparation of the NPL for consideration in subsequent updates.

Non-Contiguous Facilities

Section 104(d)(4) of CERCLA authorizes the Federal Government to treat two or more non-contiguous facilities as one for purposes of response, if such facilities are reasonably related on the basis of geography or on the basis of their potential threat to public health. welfare, or the environment. For purposes of the NPL however. EPA has decided that in most cases such sites should be scored and listed individually because the HRS scores more accurately reflect the hazards associated with a site if the site is scored individually. In other cases, however, the nature of the operation that created the sites and the nature of the probable appropriate response may indicate that two noncontiguous sites should be treated as one for purposes of listing and EPA has done so for some sites on the final NPL.

Factors relevant to such a determination include whether the two sites were part of the same operation. If so, the substances deposited and the means of disposal are likely to be similar, which may imply that a single strategy for cleanup is appropriate. In addition, potentially responsible parties would generally be the same for both sites, indicating that enforcement or cost recovery efforts could be very similar for both sites. Another factor is whether contamination from the two sites are threatening the same ground water or

surface water resource. Finally, EPA will also consider the distance between the non-contiguous sites and whether the target population is essentially the same or substantially everlapping for both sites, bearing in mind that the HRS uses the distance of three miles from the site as the relevant distance for determining target population.

Where the combination of these sectors indicates that two non-contiguous locations should be addressed as a single site, the locations will be listed as a single site for purposes of the NPL. While the nature of the listing may be a guide to prospective response actions, it is not determinative; EPA may decide that response efforts, after all, should be distinct and separate for the two locations. Also, EPA may decide to coordinate the response to several sites listed separately on the NPL into a single response action when it appears more cost-effective to do so.

VIL Changes From the Proposed NPL

The Agency received a total of 343 comments on 217 of the sites listed on the proposed NPL. General comments on the NPL are addressed throughout this preamble. Significant comments regarding specific sites are addressed in the Support Document for the National Priorities List, previously cited. A number of the site-specific comments addressed similar issues, and EPA's approaches to those common issues are presented in this section.

A total of 144 HRS score changes have resulted from the Agency's reviews of comments and other information, and these are summarized in Table L EPA determined that a total of five sites that had been proposed have HRS scores below 28.50 and should not be included on the NPL For seven sites, the Agency is still considering the comments. received concerning those sites and was unable to reach a final decision on listing in time for this publication. EPA will continue to evaluate these sites and make a final decision on them in a future update to the NPL. In one instance, where cleanup actions have adequately addressed the problems. EPA determined that a site should be deleted from the proposal and not included on the final NPL. In addition. two States have revised their designations of top priorities. These items are addressed below.

Waste Quantity. A number of commenters said that the waste quantity values essigned under the HRS were too high because EPA had included the non-hazardous constituents of the hazardous substances in calculating the quantity of waste located at the facility. This issue was raised and resolved

when the Agency adopted the HRS. In the presimile to that publication (47 FR 31100 July 16, 1982), SPA addressed the rationale for including all constituents. including the non-hazardous portions of the materials, in the calculation of the quantity of hazardous wester at a site. Briefly stated, the rationale for the Agency's approach is that detailed informetion of the portion of the total substances at a site that consist of bezardous constituents is expensive to determine, and therefore, because of the need to nee a consistent method of evaluation of this factor at many sites nationwide cannot be required as an element increasing for HRS scoring. EPA recognizes that most hezardous wastes contain some fractions of non-hazardous substances, and this fact was taken into eccount when the rating scales for waste quantity were established. In most metancies a very small amount of the hazardous anbstances can have a significant impact on public health. welfare or the environment. The Agency did not fevies waste quantity values in response to comments the non-hazardous constituents.

Consideration of Flow Gradients. In some instances commenters maintained that, based apoli their conclusions regarding prospective movement of contaminants in ground waters, the values assigned by SPA to population served by ground water are too high. The HRS, however, specifies that all the population using the aquifer of concern within a three mile radius of the facility should be included in the calculations of population served by ground water. The Agency's approach is based on the difficulty of predicting precisely the movements of ground water.
furthermore, in establishing the rating scales, the Agency took into account the fact that most walls within the three mile radius would not be affected. As was the cese with the waste quantity issue, this faste was addressed and resolved in adopting the HRS in July 1982. The retionale for the Agency's approach is further addressed in the preemble to the NOP (47 FR 31190-91. July 16, 1962) and is equally applicable

Scoring on the Brisis of Current Conditions. Bome commenters felt that EPA should take current conditions into account which scoring sites where response actions have reduced the hazards posed by the site, EPA scored sites for inclusion in the fifth based on the hazards that existed before any response actions were initiated. This policy was explained in the preamble to the final revisions to the NCP (47 FR 31187. July 16, 1982). The Agency

explained that public agencies might have been discouraged from taking response if such actions could lower HRS score and prevent a site from ber included on the NPL This has turned to be the case, as at least one State some EPA Regional Offices have actually sought reassurances prior taking emergency action at sites that site's HRS score would not be lowered as a result of the response action. Alternatively, some private parties 😤 might have only taken action suffic to lower the score to the point that it would not be listed on the NPL but would not be completely cleaned up \ Those types of score manipulations could be accomplished by such action as temporarily removing wells from service to lower target scores, or removing wastes from a site to lower 2 waste quantity scores while failing to address contaminated ground waters by remedying only air discharges when ground or surface water contamination also present a problem. Therefore, EPA was and is concerned that scoring on the basis of the latest conditions at a site could encourage incomplete solutions that might leave significant health threats unaddressed.

Even where the response actions occurred before the listing process began, EPA believes that these actions should not be considered when scoring the site for the NPL. The ability of the HRS to approximate risk at a given sile is based on a number of presumed relationships between the various factors considered in calculating the HRS scores. When partial response actions are conducted, the validity of these relationships for the purpose of : approximating the risk posed by a site may be affected. For this reason, if the site is rescored taking the response actions into account, the drop in score that may result might not reflect a commensurate reduction in the level of risk presented by a site.

For example, the factor of hazardous waste quantity, when considered with other factors that predict the toxicity of the substances and the likelihood of release, helps predict how extensive the harm from a release can be. For a site that has been in existence for some time, however, hazardous substances may already have begun migration toward ground water or surface water. the hazardous materials on the surface are then removed, and the site is scored according to conditions existing after removal, the site would be assigned ... negligible value for waste quantity. though substantial amounts of the material may still be under the site and a potential threat to the public health.

her example is where some of the i population at risk has been ed with alternative drinking water a. In such a case, the population factor might be resoured quite en where the alternative supplies aporary, costly, or limited in . In addition, rescoring in this on could penalize residents for ne alternative supplies by ng the priority of the site or us it from the list and thereby ding completion of proper ial actions. A final reason is that use action at sites is an ongoing sa, and it may become unduly. asome to continually recalculate ; to reflect such actions. ere response actions have already initiated by private parties or er agency. Histing such sites will e EPA to evaluate the need for a complete response. Inclusion on PL therefore does not reflect a nent that responsible parties are g to address the problems. The cy believes, therefore, that this sach is appropriate, and consistent the purpose of the NPL as stated in gislative history of CERCLA.

Small Observed Release. Some commenters maintained that EPA incorrectly assigned values for observed releases to ground waters because the measured concentrations of the substances involved were below the regulatory limits specified under the Safe Drinking Water Act. The ERS states:

If a contaminant is measured (rejardless of frequency) in ground water or in a well in the vicinity of a facility at a significantly fin terms of demonstrating that a release has occurred, not in terms of potential effects) higher level than the bedignound level, then a release has been observed (NCP).

Appendix A. § 3.1, 47 FR 21224, July 26, 1963).

This scoring instruction is based on the fact that the observed release factor is considered for purpose of estimating the likelihood that substances can migrate from the site. When a release is observed in any quantity, as long as the concentration is above background level, that likelihood is above prount, and this factor receives the maximum score of 45. The observed release factor is not intended to reflect the level of hazard presented by the particular release

observed. The hazard presented is. rather, approximated by the total score, incorporating the observed release factor indicating the likelihood of migration with other factors such as waste quantity, toxicity, and the persistence of the substance. These combined factors are indicative of the possibility of future releases of much higher amounts. Furthermore. concentrations of substances migrating ' in the environment tend to show extreme variation through time and space. Given that only periodic sampling is feasible in most instances, requiring contaminants to exceed certain levels before assigning an observed release could exclude many sites from the NPL which may be endangering the public. The retionale for this approach is further discussed in the preamble to the NCP (47 FR 31188 (July 18, 1982)).

Summary of Score Changes. A summary of the 144 sites where EPA's review of comments and new data resulted in a final score that changed from the score as originally proposed is shown in the table below:

BELLING CODE 6560-50-M

REFERENCE 2

CITIZEN PARTICIPATION PLAN

Garvies Point Glen Cove, New York Site No. 130032

7/17/89

Prepared By:

Fred C. Hart Associates, Inc. 530 Fifth Avenue New York, New York 10036

July 17, 1989

TABLE OF CONTENTS

<u>Section</u>		<u>Page</u>
1.0	INTRODUCTION	1
2.0	SITE HISTORY 2.1 Ownership 2.2 Dredging Activities 2.3 Landfilling Operations 2.4 Site Activities After 1983.	5 5 7 9
3.0	PROJECT DESCRIPTION. 3.1 Objectives of Remedial Investigation. 3.2 Description of RI Activities. 3.2.1 Aerial Photograph Review. 3.2.2 Preparation of Site Topographic Map. 3.2.3 Surface Water and Sediment Sampling. 3.2.4 Wetlands Sampling. 3.2.5 Air Sampling Program. 3.2.6 Shallow Test Boring Program. 3.2.7 Shallow Monitoring Well Installation. 3.2.8 Deep Test Boring Program. 3.2.9 Deep Monitoring Well Installation.	
4.0	CONTACT LISTS	14
5.0	DOCUMENT REPOSITORY	16 16 16
6.0	DESCRIPTION OF CITIZEN PARTICIPATION ACTIVITIES FOR EACH ELEMENT OF THE REMEDIAL PROGRAM	17 17 18 18
7.0	GLOSSARY OF KEY TERMS AND MAJOR PROGRAM ELEMENTS	20

LIST OF FIGURES

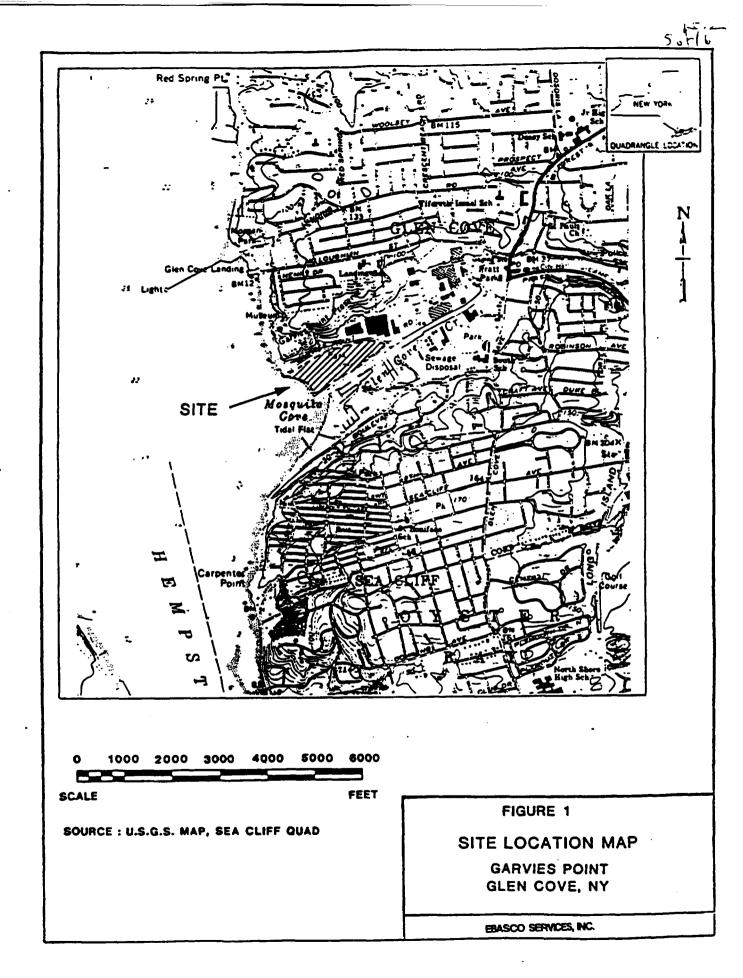
Figure No.	<u>Heading</u>	Page
1-1	Site Location Map	2
1-2	Site Map	3
2-1	Disposal Area For Materials Dredged From Glen Cove Creek in 1960 and 1965	8
i		
	LIST OF TABLES	
2-1	Garvies Point Site Ownership History	6

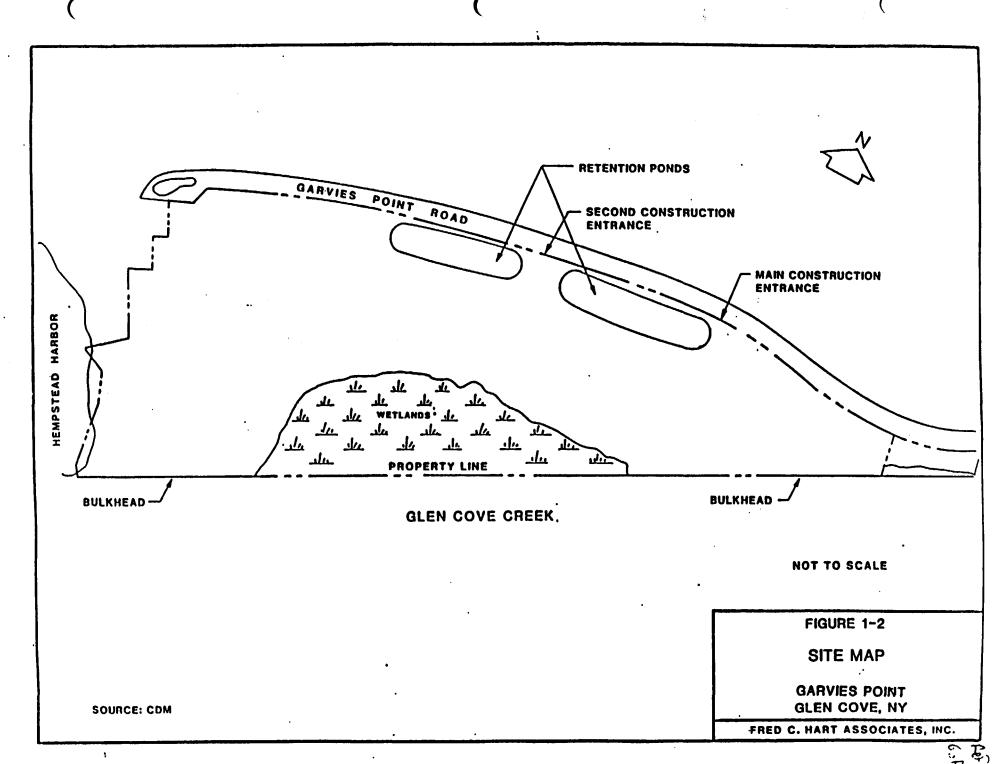
1.0 INTRODUCTION

The Garvies Point Condominium Site (hereinafter referred to as the site) is located in Glen Cove, New York at the end of Garvies Point Road. A location map of the site is presented in Figure 1-1. The site is bounded by Glen Cove Creek to the south, Hempstead Harbor to the west, the Garvies Point Preserve to the north and the Glen Cove Anglers Club to the east. A site map is presented in Figure 1-2. The total area of the site encompasses approximately 19 acres, including a section of wetlands along Glen Cove Creek.

The New York State Department of Environmental Conservation (NYSDEC) placed this site on the state's list of inactive hazardous waste disposal sites on January 7, 1986. At that time, the site (No. 130032) was assigned a rank of 2a which is a temporary classification given to sites that have inadequate and/or insufficient data for inclusion in any of the other classifications. The current owner of the site, Village Green Realty at Garvies Point, Inc. was requested by the NYSDEC in 1985 to conduct field investigations to determine if inorganic and/or organic constituents were present in different environmental media at the site. The initial test results of that investigation prompted the NYSDEC to change the classification of the site to a 2, which requires immediate action.

As a result of placement on the state's inactive hazardous waste disposal site list, Village Green Realty at Garvies Point, Inc. (Respondent) has entered into an Order on Consent with New York State Department of Environmental Conservation (NYSDEC). This Order on Consent calls for the development of a Remedial Investigation (RI) Work Plan, implementation of that Work Plan, preparation of an RI report and a subsequent scope of work for an engineering study of feasible remedial alternatives. The goals of the RI, as set forth in the order, are to determine health and environmental hazards, if any, in connection with the site; and to identify all areas of soil and water contamination at the site.





As part of the Remedial Investigation, Village Green Realty at Garvies Point, Inc. will also conduct a citizen participation program. This program will promote an understanding of the remedial activities at the site and will provide an opportunity for the collection of public information that will enable Village Green Realty at Garvies Point, Inc. to develop a comprehensive remedial program which is protective of both public health and the environment.

2.0 SITE HISTORY

2.1 Ownership

The history of site ownership was determined by searching Nassau County property records. The record search conducted by RTP Environmental Associates, Inc. (RTP) determined that recorded deeds to the property date back to 1899 when much of the site and surrounding Glen Cove Creek was assigned to Nassau County. A chronology of the owners is presented in Table 2-1. As indicated in Table 2-1, there has never been an industrial owner of the property except for The CONMAR, Inc. Group. This group purchased the property in 1979 with the intent of constructing a residuals transfer station on a portion of the eastern third of the site. The residuals were to be accepted from surface transports and transferred to barge transports docked in Glen Cove Creek. Although preliminary plans were developed, no such facility was ever constructed.

2.2 <u>Dredging Activities</u>

The United States Congress authorized the United States Army Corps of Engineers (USACE) to maintain Glen Cove Creek in 1925. Local government was to pay half of the costs and provide an acceptable disposal site for the dredge materials. The initial dredging took place from August 1933 to May 1934. The creek was dredged from Mosquito Cove in Hempstead Harbor for a length of approximately 0.7 miles upstream to a width of 100 feet and depth of 8 feet. The remaining 0.3 mile upstream portion to the head of navigation was dredged to a width of 50 feet and depth of 8 feet. A total of 195,000 cubic yards of material were removed. There are no available records regarding the disposal site for this material.

The channel has been dredged an additional three times since the initial work in 1933 and 1934 was done. In 1948, 26,500 cubic yards were removed but there are no available records to indicate where this material was disposed. In 1960, 27,100 cubic yards were dredged from the lower portion of Glen Cove Creek. According to information available from the USACE, this material was disposed of on the Garvies Point site. Finally,

(1823n-5)

TABLE 2-1

GARVIES POINT SITE OWNERSHIP HISTORY

Lot #	<u>Owner</u>	Deed Date
26 & 27	Hm. H. Seaman .	9/21/09
(later redesignate	d)	
424 & 546	E.S. Appleby, et al. City of Glen Cove Realty Assoc. John White Ridgewood Platear Realty Assoc. Glen Cove Realty Corp. J. Graham Creek Develop. Corp. I.I. Miller City of Glen Cove Nassau County City of Glen Cove I.I. Miller Lee Langbaum CONMAR Blders. James O'Connell Glen Cove Development Corp. Village Green	6/26/31 9/13/46 10/14/47 10/4/49 10/5/49 12/12/51 01/10/56 12/20/56 12/22/56 02/13/58 04/28/70 09/13/74 11/19/74 09/02/76 03/26/79 12/28/79 04/16/81 08/15/83
551 & 556	(same as 424 and 546 prior to 1 Glen Cove Urban Renewal Glen Cove Community Development Village Green	04/12/77

^{*} Deed date not in records

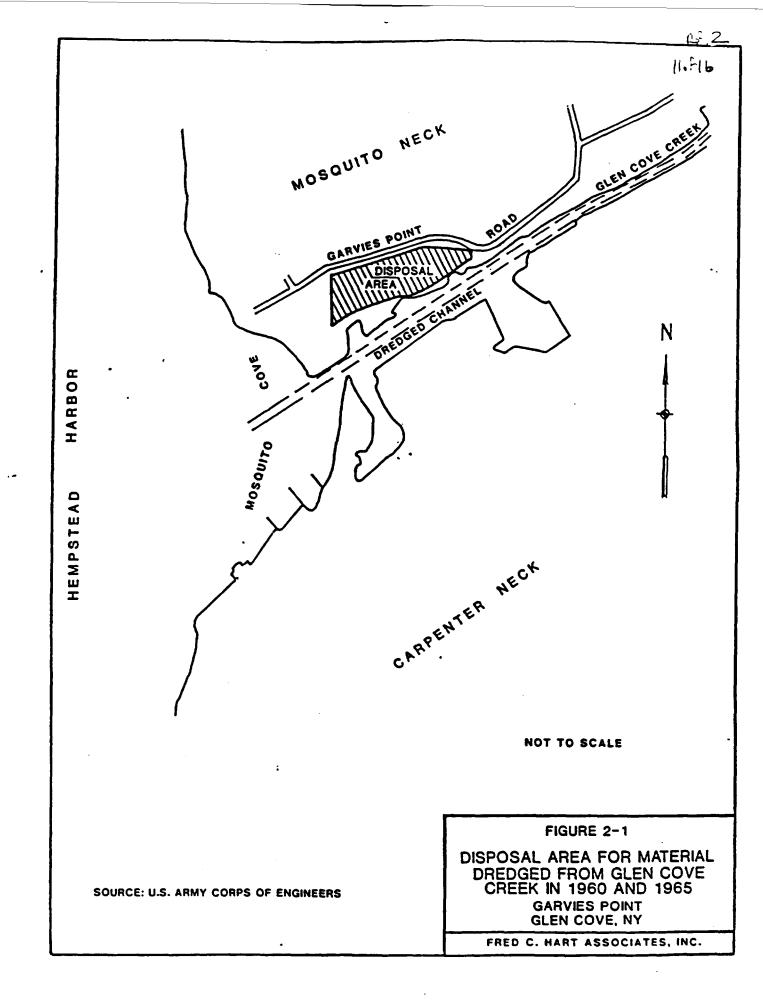
6,300 cubic yards were dredged in 1965 and reportedly also disposed of on the site. The approximate disposal area for material removed in 1960 and 1965 is shown in Figure 2-1.

An USACE proposal to dredge Glen Cove Creek in 1979 was not implemented. Certain analytical tests conducted in sediment samples from the creek indicated a potential for the creek sediments to release PCBs, iron, and cyanide in concentrations greater than was currently present in the receiving water body (Glen Cove Creek). The 1979 proposal by the USACE included disposal of the dredge materials on the site. However, the NYSDEC prohibited this land disposal option after the dredge material was deemed to be hazardous.

2.3 Landfilling Operations

According to Nassau County Department of Health (NCDOH) records, the City of Glen Cove used the site as a municipal sanitary landfill beginning in 1971. Incinerator residues, wastewater treatment plant sludges and street debris were disposed of at the site. However, use of the site as a disposal area may have begun earlier. During the early years of city ownership, the records show that complaints had been received by the county. These complaints were related to the burning of rubbish at the site and to odors allegedly coming from uncovered sewage sludge. The City of Glen Cove was responsible for the site and occasionally cleaned and removed debris from the site.

RTP reports that the landfilling activities at the site were corroborated by Mr. Donald Aitken, a former NCDOH sanitarian who was responsible for periodically inspecting this and other landfills. Mr. Aitken explained that to his knowledge, household debris was disposed of in the western section of the landfill. The primary area of landfilling occurred in the central section of the property. The garbage consisted of typical household garbage, construction debris, catch basin sediments and sludge from the City sewage treatment plant. Mr. Aitken was unsure of the exact boundary of the eastern border of the landfill. He did acknowledge that at one time a 20 foot high sand berm existed along



the northern border of the site. This berm was subsequently leveled and graded after 1983. A soil berm also existed along the southern border of the site. Mr. Aitken did not recall any incidents of industrial or potentially hazardous waste or ash disposal at the site. He also stated that the landfill was still active into the early 1980s just prior to the purchase by Village Green Realty at Garvies Point, Inc.

In NCDOH records dating back to 1973, references are made to the disposal of incinerator ash, sewage sludge, household debris and other sanitary fill on various portions of the site. Prior to about 1975, the discarded debris in the landfill was burned, apparently to reduce the volume of the discarded materials and for rodent control. There are no available records describing where these disposal operations took place. However, based on topography and the aerial photographs, it appears that the center of the current site was the primary disposal area.

2.4 Site Activities After 1983

The property was purchased by Village Green Realty at Garvies Point, Inc. in the fall of 1980. Since that time, bulkheads have been built along Glen Cove Creek and the western end of the site bordering Hempstead Harbor. The bulkheads were backfilled with clean fill. Approximately one third of the distance along Glen Cove Creek was not bulkheaded in order to preserve an estuarian habitat. Two lined retention ponds were constructed near Garvies Point Road. The purpose of the ponds is to collect surface runoff and allow solids to settle out of the water before the water is released to Glen Cove Creek. The intent of the liners in these two ponds is to prevent infiltration of stormwater into the subsurface which may be comprised of landfill materials.

Both wooden and concrete piles have been driven into the subsurface over much of the site. The purpose of these piles is to provide structural integrity to residential units planned for the site. There are currently two residential units whose frames have been constructed in the eastern portion of the site. The construction of these units was suspended by the developer. These framed residential units include

elements of a gas collection system which had been had specified prior to construction. Finally, a stockade and chain link fence exists along the northern and eastern site boundaries.

3.0 PROJECT DESCRIPTION

3.1 Objectives of Remedial Investigation

The property was purchased by Village Green Realty at Garvies Point, Inc. in 1980 with the intention of developing a residential complex on the site. Village Green Realty at Garvies Point, Inc. in the spirit of cooperation and without admitting liability for the disposal of industrial or hazardous waste at the site, has consented to enter into and carry out the elements of the Order on Consent with New York State Department of Environmental Conservation (NYSDEC). The goals of the RI, as set forth in the order, are to determine health and environmental hazards, if any, in connection with the site; and to identify all areas of soil and water contamination at the site.

In addition to the RI, a radiological survey of the site will be conducted. This survey is described in a separate work plan and will be conducted prior to the start of the RI field activities. The goals of the survey are to assess the potential hazard from radioactive materials, if any, deposited at the site by local industries. The amount of radiation above the ground surface will be measured with hand held radiation detecting instruments. If above background readings are measured on the instruments, up to ten samples will be collected for laboratory analysis.

3.2 Description of RI Activities

- 3.2.1 <u>Aerial Photograph Review</u>. HART will review aerial photographs taken of the site from 1950 through 1986 to define the locations of potential dredge and/or landfill materials and to observe any topographical changes at the site.
- 3.2.2 <u>Preparation of Site Topographic Map</u>. Since there is no topographic or scaled base map available for the site, a survey company will be subcontracted to prepare such a map. The map will be constructed on a scale of one inch equal to 100 feet at an appropriate contour interval.

(1823n-11)

- 3.2.3 <u>Surface Water and Sediment Sampling</u>. In order to characterize the surface water and sediment at Garvies Point, samples will be collected from the two retention ponds and the pipe that discharges from these ponds to the creek. The ponds collect surface water and sediment drainage from the entire site and are therefore representative of the site as a whole. In addition to these three sample locations, two surface water and two sediment samples will be collected from seeps noted on the southeastern slope of the site, adjacent to the wetlands area. These samples will be analyzed for Target Compound List (TCL) and Target Analyte List (TAL) constituents. These samples will provide the most accurate indication of the concentration of any compounds leaving the site and entering the creek.
- 3.2.4 <u>Wetlands Sampling</u>. In order to evaluate any potential impact of the site on the adjacent wetlands, five sediment samples will be collected from the wetlands. The samples will be collected during low tide when the maximum amount of wetlands area is exposed. These samples will be analyzed for TCL organics and the inorganics.
- 3.2.5 Air Sampling Program. Two different air sampling techniques will be used to determine the presence of any subsurface soil gas at the site. The first method will entail the collection of vapor samples from shallow holes in the ground approximately 3 feet deep and 3/8 inch in diameter. These vapor samples will be screened in the field to determine locations where vapor samples will be collected for laboratory analysis. Collection of the second set of soil vapor samples will require inserting a hollow probe approximately two feet into the ground and then pumping soil vapor through the probe and into sample tubes. Approximately eight (8) samples will be collected for laboratory analysis. Ambient air (upwind and on-site) will also be analyzed during this program. The laboratory results will:permit calculation of surface emission rates for any volatile organic compounds detected.
- 3.2.6 <u>Shallow Test Boring Program</u>. A total of 13 shallow borings will be drilled at the site to gather additional data regarding the nature of potential organic and inorganic compounds in the soil fill area. These

borings will also provide lithological information to delineate the lateral continuity of the clay layer at the site. Up to ten samples will be collected from each boring and these samples will be tested in the field for pH, conductivity and volatile organics to determine which should be submitted for laboratory analysis. Up to nineteen samples will be selected from the borings for analysis.

- 3.2.7 <u>Shallow Monitoring Well Installation</u>. Up to six shallow monitoring wells will be installed at the site, which in combination with the existing four wells at the site will provide the necessary data to define the vertical extent of potential groundwater contamination at the site. The monitoring program will also focus on characterizing upgradient or off-site organic and/or inorganic contamination. Groundwater samples will be collected from all ten shallow wells and submitted for laboratory analysis.
- 3.2.8 <u>Deep Test Boring Program.</u> The objective of the deep boring program is to evaluate more fully the connection between the fill material above the clay layer and the fill material found along the bulkhead on the southeastern, and possibly the southwestern, side of the site. In addition, an attempt will be made to determine the thickness of the clay layer along the northwestern side of the site. Soil samples of the fill material will also be collected from two of the three borings for TCL and TAL analysis. Depending upon the results of the boring program, up to three of these borings may be finished as monitoring wells.
- 3.2.9 <u>Deep Monitoring Well Installation</u>. Depending upon the results of the deep test boring program, one of several objectives may be accomplished by installing deep monitoring wells at the site. If all three wells are installed, both horizontal and vertical hydraulic gradients at the site may be assessed as well as the quality of deeper water bearing zones. If no water bearing zone is found at the upgradient well location within fifty feet of the surface and only two wells are installed, only vertical hydraulic gradients at the site may be assessed. The two deep wells will be sampled to assess groundwater quality at depth in the fill material near the bulkhead. However, the data will have to be interpreted carefully since no upgradient data will be available.

REFERENCE 3

*** DRAFT

REMEDIAL INVESTIGATION

WORK PLAN

GARVIES POINT CONDOMINIUMS

GARVIES POINT ROAD

GLEN COVE, NASSAU COUNTY

MARCH 1988

Prepared By
RTP Environmental Associates, Inc.
and
Fanning, Phillips, and Molnar

DRAFT

*** DRAFT ***

REMEDIAL INVESTIGATION WORK PLAN

TABLE OF CONTENTS

ECTION	TITLE	PAGE #
1.0	INTRODUCTION ·	
2.0	OBJECTIVE	
3.0	HISTORICAL INFORMATION SUMMARY 3.1 Site Reconnaissance 3.2 Previous Site Studies 3.3 Aerial Photography 3.4 Site Map 3.5 Regional Hydrology, Hydrogeology and Climatology 3.6 Public and Local Water Supply Wells 3.7 Previous Dredging Activity 3.8 Landfilling Operations 3.9 Site Activities After 1983	
4.0	SAMPLING PLAN	
5.0	BOREHOLES AND MONITORING WELL INSTALLATION 5.1 Borehole Drilling 5.2 Well Installation	
6.0	SAMPLING AND ANALYSES 6.1 Volatile Fraction of the HSL Test Boring 6.2 Property Line Monitoring Well Installation 6.3 Downgradient and Deep Monitoring Well Installation 6.4 Site Interior Test Boring Installation 6.5 Site Geohydrology 6.6 Glen Cove Creek Characterization 6.7 Groundwater Sampling 6.8 Decontamination 6.9 Sampling Equipment 6.10 Sample Containers 6.11 Air Quality Sampling 6.12 Soils Sampling 6.13 Creek Sampling	

TABLE OF CONTENTS (PAGE 2)

SECTION	TITLE	PAGE #
7.0	QUALITY ASSURANCE AND QUALITY CONTROL 7.1 Samples Collected for Quality Control Purposes 7.2 Duplicate Samples 7.3 Travel Blanks 7.4 Field Blanks 7.5 Intralaboratory Splits 7.6 Interlaboratory Splits	
8.0	HEALTH & SAFETY PLAN	
9.0	DATA REPORTING AND ANALYSIS	
APPENDIX A:	DATA SUMMARY REPORT	
APPENDIX B:	CDM, 1986 REPORT EXCERPTS LKB, 1985 REPORT EXCERPTS	
APPENDIX C:	PROPOSED 1979 GLEN COVE CREEK DREDGING CORRESPONDENCE	

1.0 INTRODUCTION

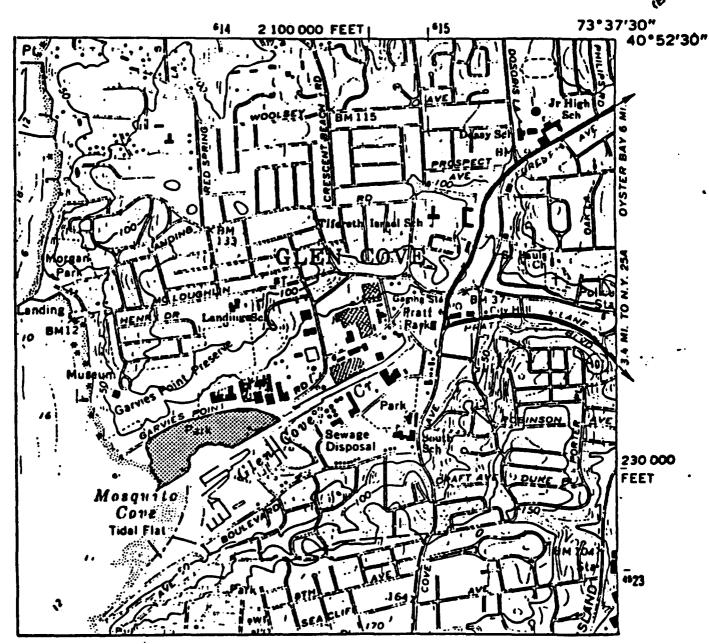
The Garvies Point Condominium Site is located in Glen Cove, New York at the end of Garvies Point Road. The site is bounded by Glen Cove Creek to the south, Hempstead Harbor to the west, the Garvies Point Preserve to the north and the Glen Cove Anglers Club Marina to the east. The site is shown on Figures 1.1 and 1.2 As outlined in Figure 1.2, the site consists of approximately 19 acres, which includes a section wetlands along Glen Cove Creek.

The current site owner, Village Green Realty at Garvies Point Inc. has entered into a Order On Consent on the Garvies Point Condominium Site with the New York State Department of Environmental Conservation (NYSDEC) under Article 27, Title 13 of the Environmental Conservation Law (ECL) of the State of New York. Formerly the site was owned by the City of Glen Cove and a portion of the site was used as a landfill by the City of Glen Cove.

In 1985, the site owner by request of the NYSDEC performed a preliminary site investigation to determine if hazardous wastes were located on site (CDM, 1986). The site investigation determined that hazardous materials were present and the NYSDEC then reclassified the site as an inactive hazardous waste disposal site as that term is defined in Section 27-1301 (2) of the ECL. The site number designation under the ECL is No: 130032. Furthermore, the NYSDEC stated in the Order On Consent that the hazardous and industrial substances, hazardous waste constituents and toxic degradation products thereof, at and in the vicinity of the site constitute a significant threat to the environment. Pursuant to ECL 27-1313 (3)(a) the Commissioner of Environmental Conservation may order the owner of such a site and/or any person responsible for the disposal of hazardous wastes at such site (1) to develop an inactive hazardous waste disposal site remedial program and (2) to complete such a program within reasonable time limits.

SEA CLIFF QUADRANGLE NEW YORK 7.5 MINUTE SERIES (TOPOGRAPHIC)

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GARVIES POINT SITE

FIGURE 1.1: PROJECT SITE

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eff	ective			_ •							

The goals set forth in the Order On Consent are: (1) determine the health and environmental hazards and potential hazards in connection with the site and (2) identify all areas of soil and water contamination at the site.

In response to the Commissioner's Order, Village Green Realty has developed this work plan to accomplish the goals stated above.

This work plan is comprised of several sections. In summary the plan consists of a site reconnaissance, sampling and analysis plan, investigation protocols, quality assurance and control, health and safety procedures and reporting protocol. In preparing and completing this work plan, Village Green Realty, its consultants and assigns do not admit liability for the disposal of industrial or hazardous substances at the site. Such liability must be the acknowledged responsibility of the former owners of the site and any claims for damages or otherwise, therefore, are their responsibility.

2.0 OBJECTIVE

The objective of the remedial investigation is to provide additional data on site characteristics including the characteristic, location, quantity and quality of any hazardous materials on-site and the potential for on-site materials to enter air, soil and water media pathways. These data will be used in the feasibility study for the site to determine the measures necessary to fully and safely remediate any potential hazards found.

The ultimate objective of the proposed remedial investigation (RI) is to provide information on the nature and extent of materials on-site so that an effective remediation program can be implemented. The completion of the remediation will allow the issuance of a clean bill of health for the site. Subsequent to the issuance of the clean bill of health, the applicant fully intends to develop the site as a residential condominium development.

3.0 HISTORICAL INFORMATION SUMMARY

3.1 Site Reconnaissance

Site reconnaissance has been conducted by several groups including the NYSDEC. The previous site investigations have also been conducted as noted in Appendix A. Because of the nature of area, its former use as a municipal landfill and dredge spoil disposal area, and the significant amount of alteration that has occurred subsequently, the approach taken in this work plan will be to cover the area with additional observation wells, soil borings and air samples to fully characterize the site and to some extent the adjoining area of Glen Cove Creek.

Aerial photographs showing the local land use, adjoining sites, the City of Glen Cove Codisposal Plant and other industrial sites are included in Appendix A. A detailed site reconnaissance map will be developed and discussed for inclusion into the RI document.

3.2 Previous Site Studies

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Lockwood, Kessler and Bartlett (LKB) (LKB, 1985) have prepared an engineering report on the site. The report contains data on soil profiles across the site as well as other information on-site geophysics. Relevant sections of the LKB report are included as Appendix B. Data are also presented on the test borings done on-site.

A Camp Dresser & McKee (CDM) (CDM, 1986) report provides data on several groundwater wells, soil profiles, metal concentrations and other site characterization data. These data have been used in developing the proposed supplementary sampling and analysis plan presented in the following section. Relevant sections of the CDM report are included in Appendix B.

The Camp, Dresser and McKee report (CDM, 1986) discussed the results of samples from twenty shallow soil borings (2 foot deep), four deep borings (15 foot deep) and four groundwater monitoring wells. Figure 3.1 from the CDM report (attached for reference) shows the locations of the twenty shallow soil samples, the four 15 foot boreholes and the four observation wells.

The CDM shallow soil samples were analyzed for HSL-CLP metals. The composite soil samples from the 15 foot deep borings were analyzed for HSL inorganics, pesticides, PCB's and cyanide. The four soil observation well samples were analyzed for HSL-CLP volatiles, base neutral and acid extractables and inorganics.

The results of each CDM shallow soil sample were averaged together to develop an average concentration for each metal to provide a basis for identifying the highest values. Six sampling locations contained metals whose values were elevated. These six locations, are in two clusters, one cluster contains CDM samples SO4 and SO5 and the other cluster contains CDM sample SO2, S10, S11 and S12.

The results of the four 15 foot deep borings show many metal concentrations at or below the detection limits. Two of these borings showed elevated metal concentrations in the top soil layer, in the zero to six foot composite (CDM#B3 and CDM#B4). These have the most number of metals exceeding the shallow soil average. Borings 3 and 4 are in the cluster area of CDM shallow soil samples SO2, S10, S11 and S12 where higher than average metals were detected. Cyanide was not detected in the deep borehole samples at any depth.

Aroclor 1248 was present but below detection limits in all but two CDM deep borehole samples. The highest Aroclor level was detected at borehole CDM#B1 at 7.5 - 9 feet. Some pesticide compounds were present below detection

FIGURE 3.1 - CDM SAMPLE LOCATION MAP

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limits including 4-4' DDD and chlordane. Although relatively low, the highest pesticide concentration was for chlordane at borehole CDM#B1 at the 7.5 -9 foot depth. These low levels should be further delineated.

The results of the CDM aqueous analysis show volatile organic compounds present at CDM Well #2. The other three CDM wells contained concentrations of volatile organics below the detection limits or below values which were found in blank samples. CDM Well #4 contains detectable concentrations of benzene, phenol and bis (2-ethylhexyl) phthalate. An analysis of the groundwater samples also showed unidentified compounds in detectable unquantifiable concentrations (predominantly at CDM Well #4). The CDM report stated that the high levels of organics at CDM Well #2 may be due to off site sources based upon the well's location and measured water gradients.

The highest aqueous metal concentrations were found in CDM Well #4. However, all CDM water samples were unfiltered and were preserved in the field prior to transport to the laboratory for analysis. This procedure dissolves all metals in particulate form and represents unrealistically high values for true dissolved metals in the groundwater. No cyanide was detected in the CDM aqueous samples.

In summary, the CDM shallow soil analysis identified two areas of high metal concentrations. The deep boring samples identified PCB and pesticide compounds at low levels at CDM Well #2. Metal compounds were identified in all aqueous samples with the highest levels at CDM Well #4. Unidentified base neutral extractable compounds were detected in the aqueous samples. Phenol and bis (2-ethylhexyl) phthalate were detected at CDM Well #4. Cyanide was not detected in the soil or the groundwater on site.

The proposed sampling plan, as discussed in the following Section 4.0, will address and expand upon the CDM report. Compounds identified by CDM's soil sampling analysis will be included in the proposed soil sampling analysis. These compounds include HSL-metals, pesticides, base neutral and acid extractables; PCB's, phenols and cyanide. Selected samples will be analyzed for HSL volatiles. Aqueous parameters identified in the CDM report and included in the proposed sampling plan include HSL-metals, base neutral and acid extractable volatile fraction, phenols and cyanide. Even though cyanide was not detected on site, it is included in the proposed sampling plan because of the history of the dredge spoil disposal on site as discussed in Section 3.7.

3.3 Aerial Photography

As described in Appendix A, aerial photographs for the site are available from 1950 through 1986. Photo reprints of the available photography will be provided in the RI document. A review of the aerial photographs is provided in Section 3.7. The aerials indicate areas of filling that had occurred on-site during the various periods noted up to 1966, the period covering the last known deposition of dredge spoils from Glen Cove Creek on the site. A complete analysis of existing photographs will also be provided in the RI.

3.4 Site Map

A site map and plot plan of the Garvies Point Condominium Site will be developed through ground survey methods and will include significant surface, topographic and structural features and the establishment of an on-, site benchmark.

3.5 Regional Hydrology, Hydrogeology and Climatology

The regional hydrogeology and geology will be presented referencing the appropriate USGS and other relevant data. Area drainage basins and patterns including surface water hydrology will be addressed. Tidal effects and water table fluctuations at the site will be referenced. Local climatic conditions and meteorological factors including precipitation and their effects upon the site will be compiled. Data from local National Weather Service stations will be used in the RI analysis.

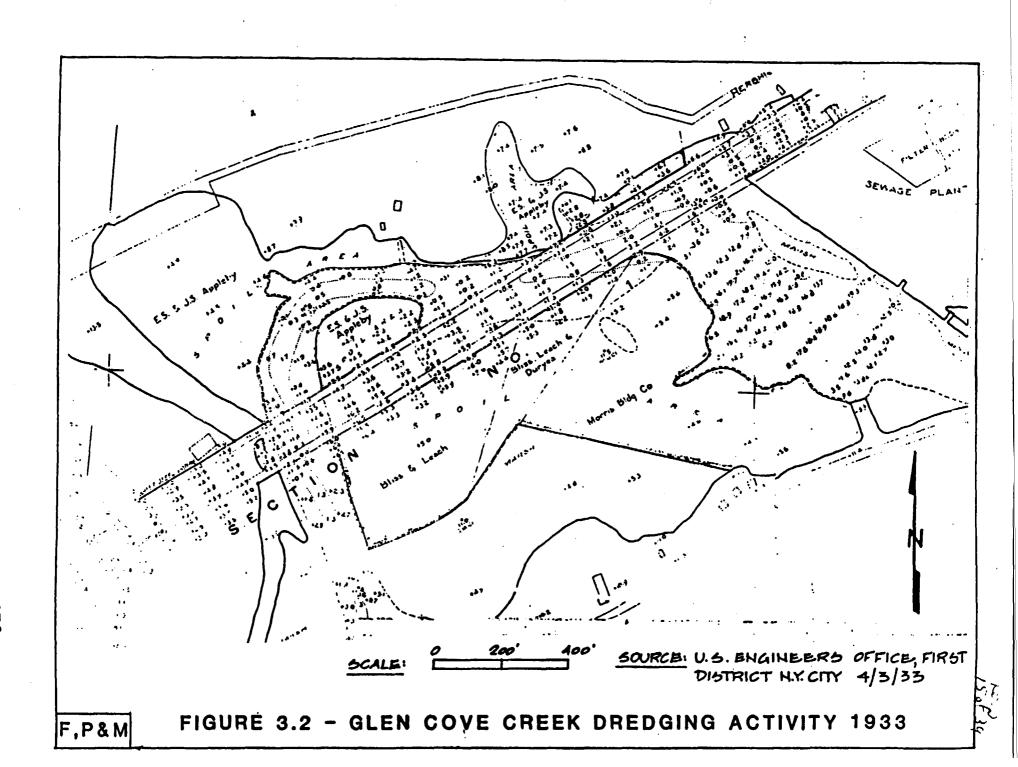
3.6 Public and Local Water Supply Wells

All potable and water supply wells within a one mile radius of the site will be mapped on a scaled base map and presented in tabular form.

3.7 Previous Dredging Activity ·

Historic records of the dredging and spoil disposal activities at Glen Cove Creek were reviewed at the Army Corps of Engineers, Navigation Section, New York District Office in New York City. Mr. S. Lew of the Navigation Section provided the files. The primary concern was the disposal of dredge spoils on the site property, the origin of the dredge spoil and their characteristics.

Based upon the existing records, dredging of Glen Cove Creek occurred in 1948, 1960 and 1965. An April 1933 map of Glen Cove Creek (Figure 3.2) obtained from the U.S. Engineers Office, First District New York City shows dredge spoil disposal areas which implies pre-1948 dredging activity. Although unconfirmed, there is a strong likelihood that such dredging did occur since the Army Corps significantly modified the original course of Glen Cove Creek from the trajectory shown in Figure 3.2 to the current straight line trajectory.



In the 1948 records of dredging activity, it was proposed that the volume of dredge material to be removed was 29,500 cubic yards. No maps of the actual dredging or spoil disposal areas were in the file. Estimates of the disposal areas were approximated from the 1950 aerial photographs.

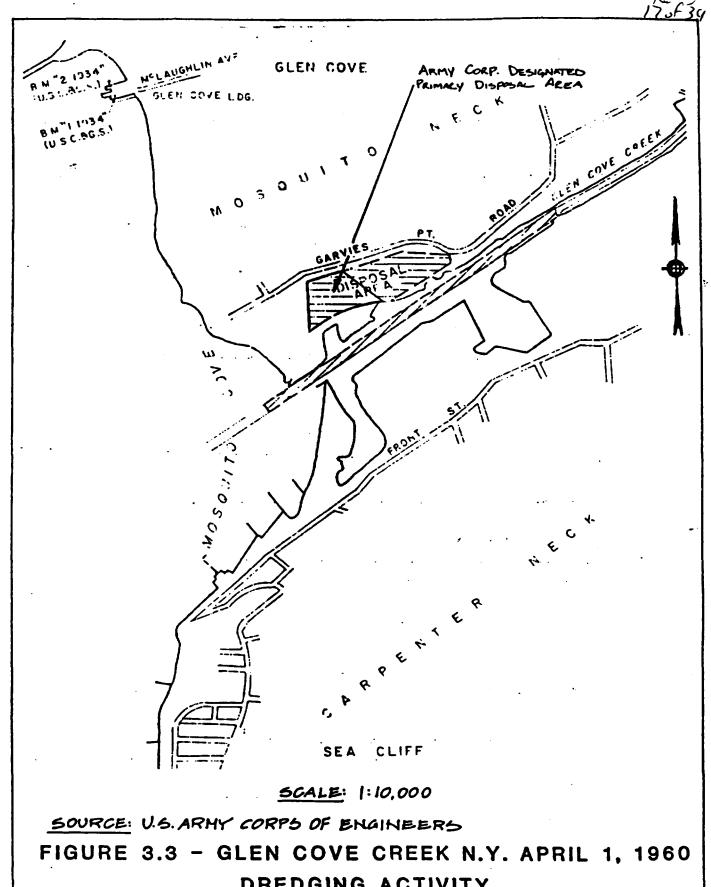
A review of the 1960 records revealed that the proposed volume of dredge material to be removed was 27,600 cubic yards.

The area to be dredged and the dredge spoil disposal area from the U.S. Army Corps of Engineers dated 4/1/60 are shown in Figures 3.3. The central and eastern sections of the property are designated as the disposal areas.

A review of the 1965 Army Corps records shows that dredging took place and the proposed dredge spoil disposal area was in the center of the site, Figure 3.4. The area to be dredged was approximately where Glen Cove Creek discharges into the Harbor.

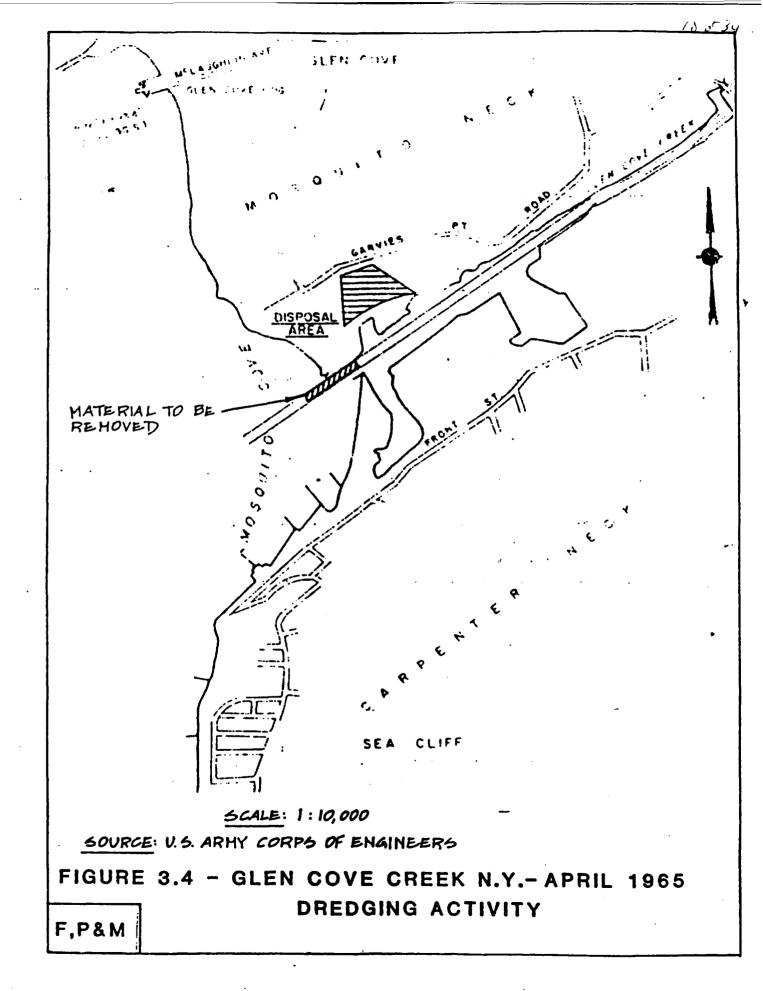
The proposed 1979 dredging activity shown in Figure 3.5 did not occur. The reasons given were that the upstream area of the creek was not properly bulkheaded for dredging activity. Additionally, a sediment sample of the creek obtained by the EPA showed levels of cyanide above regulatory guidelines. It must be assumed the sample showing the contamination was taken from the area to be dredged, although no specific location or number of samples was provided. The NYSDEC subsequently declared the proposed dredge spoils as a hazardous waste and prohibited disposal of the material in the landfill at the Garvies Point Site. Alternative disposal sites were investigated, however, dredging did not occur because other disposal areas were cost prohibitive (Appendix C).

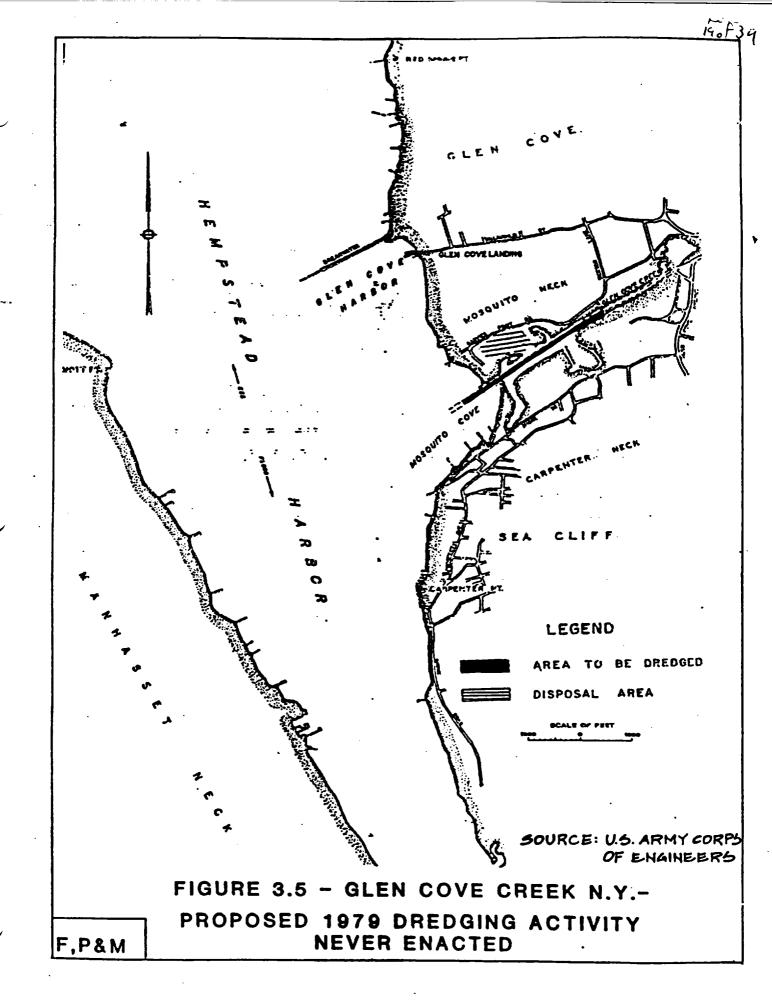
A composite sketch of the dredging activity and placement on the site has been provided in Figure 3.6.



DREDGING ACTIVITY

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3.8 Landfilling Operations

The City of Glen Cove during its ownership of the site, had used it as a landfill. Nassau County Department of Health (NCDOH) indicates that both incinerator residues and wastewater treatment plant sludges were deposited at the site dating back to 1971. The records also indicate that street debris were disposed of at the site.

These findings were also corroborated by Mr. Donald Aitken, a former NCDOH sanitarian who was responsible for periodically inspecting the landfilling activities at the site. Mr. Aitken explained that to his knowledge, household debris was disposed in the western section of the landfill. The primary area of landfilling occurred in the central section of the property involving trenching with a backhoe, filling the trench with garbage and moving to an adjacent area to repeat the process. To his knowledge, the garbage consisted of typical household garbage, construction debris, catch basin clean out material and occasionally sludge from the City sewage treatment plant. The material was not sorted.

Mr. Aitken was unsure of the exact demarcation of the eastern border of the landfill. He did acknowledge that a 20' high sand berm existed along the northern border of the site. This berm was subsequently leveled and graded after 1983. A soil berm also existed along the southern border of the site.

Mr. Aitken did not recall any disposal incidents of industrial or potentially hazardous waste or ash. He also stated, to his knowledge, that the landfill was still active into the early 1980's just prior to the purchase by Village Green Realty. In NCDOH records dating back to 1973, references are made to the disposal of incinerator ash, sewage sludge, household debris and other fill on various portions of the site. Prior to about 1975, the discarded debris in the landfill was burned apparently to reduce the volume and for rodent control. No good records were available where these disposal

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operations took place, however, based on topography and the aerial photographs, it appears that the center of the current site was the primary disposal area.

3.9 Site Activities After 1983

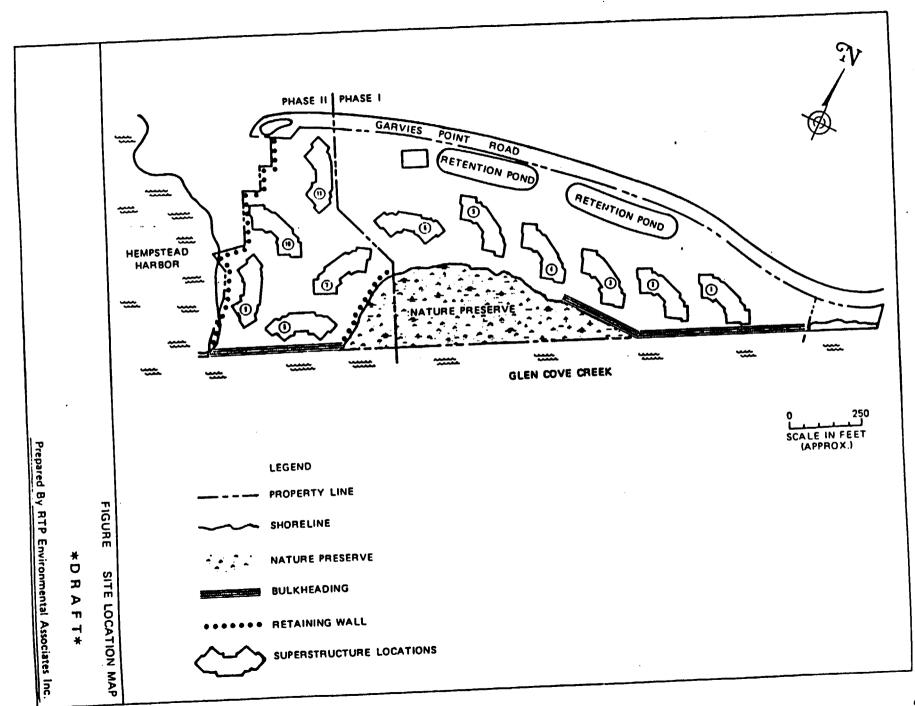
The various parcels comprising the site were officially purchased by Village Green Realty in the fall of 1983. The ownership of the site prior to 1983 has been outlined in Appendix A.

After the site was purchased by Village Green Realty, Inc., a master plan for development was prepared and approved by various City and County agencies, the NYSDEC, and the Army Corps of Engineers (with respect to the preservation of on site wetlands). The site development plan called for extensive bulkheading along Glen Cove Creek and along the western portions of the site. Many thousands of yards of clean fill were brought onto the site to fill in behind the bulkheading and well as several lower lying areas. As part of the development plan, two retention ponds were developed along the northern border of the property as shown in Figure 3.7. The intended purpose of the retention ponds was to collect surface runoff from the various portions of the site to prevent infiltration of precipitation into the old landfilled areas.

Bulkheading was initially proposed along the entire southern portion of the site. However, because of wetland concerns, approximately a third of the bulkheading was not installed to provide for the protection of the estuarian habitat along Glen Cove Creek.

The initial development plan called for multi-story structures with apartments on the ground floor. Wooden piles were driven into the areas beneath the proposed structures. Because of concerns over methane gas

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released from the underlying marsh areas, dredge spoils, and the other decomposing landfill materials, a modified plan for condominium development included several special features to prevent the migration of methane into enclosed spaces or into the living units. Additional piles were driven to support the redesigned units bringing the total number of piles driven into the site to approximately 4,000. The site development plan was approved by all regulatory agencies and construction of the superstructures was initiated. Due to financial difficulties and an injunction against the construction, only two superstructures were constructed.

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The two super structures were never completed and continue to stand on the eastern portion of the site. Wooden and concrete piles have been placed and continue to remain in over two-thirds of the site. The only section of the site that is free from structures is the western most portion. A sales pavilion was constructed on the western third of the site and a blacktop roadway was constructed to allow access to the sales pavilion. Landscaping was provided along the access roadway as well as surrounding the sales pavilion. To limit access to the site, a six foot high stockade fence and chain link fence was placed along the northern and eastern boundaries of the site.

APPENDIX A

DATA SUMMARY REPORT



RTP ENVIRONMENTAL ASSOCIATES INC.

AIR . WATER . SOLID WASTE CONSULTANTS

400 Past Avenue. Westbury. New York 11590

(516)333-4526

March 1, 1988

Mr. Robert Becherer
Regional Engineer
Division of Solid & Hazardous Waste
New York State Department of Environmental Conservation
Building 40
State University of New York
Stony Brook, New York 11794

RE: Submittal of Data on Garvies Point Condominiums

Dear Mr. Becherer:

On behalf of Village Green Realty at Garvies Point, Inc., the following data report has been compiled to satisfy the requirements stipulated in the NYSDEC Order On Consent for the referenced site.

The Order requires a brief site history, a description of previous investigations, a historical inventory of serial photography and other data not previously provided to the NYSDEC. The following report summarizes our findings.

Please review the information and contact me if you have any questions or comments or know of additional information that may be helpful for this site investigation.

Sincerely yours,

RTP ENVIRONMENTAL ASSOCIATES, INC.

Kenneth J. Skipka

Principal

KJS/erl

cc: N. Nyman

D. Rothberg, Esq.

R. Piaggione, Esq. (w/o attach)

K. Phillips, PhD

D. Elias

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DATA SUMMARY REPORT GARVIES POINT CONDOMINIUM SITE, GLEN COVE, N.Y.

1.0 INTRODUCTION

An Order On Consent exists between Village Green Realty at Garvies Point, Inc. and the State of New York: Department of Environmental Conservation regarding the Garvies Point Condominium Site in Glen Cove, New York (hereinafter known as the site). Village Green Realty is to provide the State with "all data within its possession or control regarding environmental conditions on-site and off-site, to the extent that such data has not heretofore been provided to the Department".

At a minimum, these data shall include:

- a. A brief site history,
- b. A description of the results of all previous investigations, and
- c. A historical inventory and best available copies of all aerial photography available for the site.

This document provides the above data directly, summarizes the studies that have been conducted, and gives the appropriate references for securing the data/information identified above.

2.0 SITE OWNERSHIP HISTORY AND USES

In order to determine former site ownership, a search of the Nassau County property records was performed. The search, in general, determined that the recorded deeds to the property originally dated back to 1899 at which time the site and much of the area surrounding Glen Cove Creek was assigned to Nassau County. In 1909, Wm. H. Seaman acquired ownership. In 1931, the

City of Glen Cove was deeded the property from the Appleby's. (There is a gap in the records regarding how the Appleby's acquired the property.)

After 1931, many changes in ownership of the property occur. Table 1 lists the owners and when the deed to the property was conveyed. A map of the site is provided as Figure 1.

As shown in Table 1, there has never been an industrial owner of the property except for the CONMAR, Inc. Group. The CONMAR Group purchased the property with an intent of constructing a residuals transfer station on approximately the eastern third of the site. The residuals were to be accepted from surface carriers and transferred to barge transports docked in Glen Cove Creek. Although preliminary plans were developed in 1980, no such facility was ever constructed.

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Village Green Realty acquired the several lots that comprise the property during the period 8/15/83 to 10/4/83.

The Nassau County Department of Health (NCDOH) was contacted and their files were reviewed to determine the general uses of the property. Available Nassau County records begin in 1963 at which time the site was being considered by the City of Glen Cove for the landfilling of City incinerator ash and residuals and sewage sludge from the City wastewater treatment plant. Records show that the site was actively used as a disposal site for incineration residuals and for sewage sludge. Nassau County records also indicate that the site was also used by unauthorized individuals for disposal of rubbish. In this case, the majority of the rubbish disposal was apparently confined to household debris. The City of Glen Cove was also accused of illegally using the area to dispose of City street debris. During the early years of City ownership, the records show that complaints had been received by the County. These complaints related to the burning of

TABLE 1

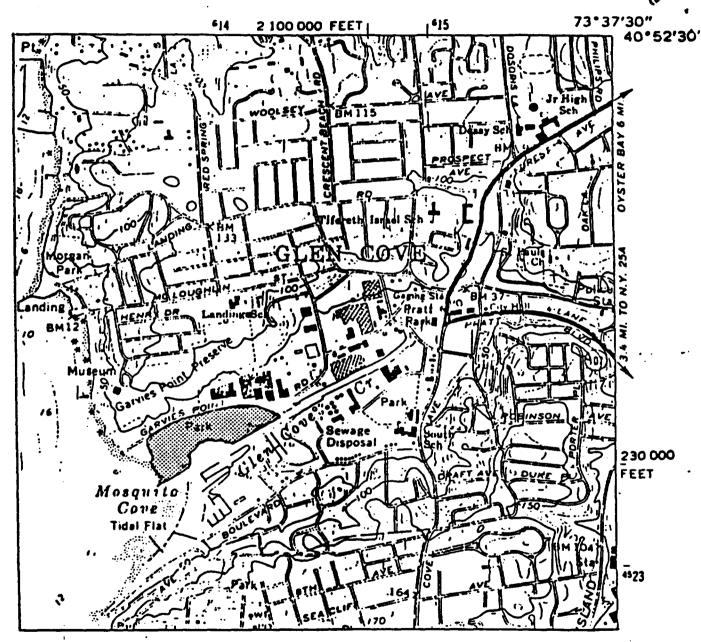
GARVIES POINT SITE OWNERSHIP HISTORY

The site previously consisted of several parcels on Nassau County Tax Maps in Section 21 Block A, Lots 551, 514, 546, 555 and 556. The ownership according to the deeds filed with Nassau County:

Lot #	Owner	Deed Date		
26 & 27	Wm. H. Seaman	9/21/09		
(later redesignated))			
424 & 546	E.S. Appleby, et. al.			
	City of Glen Cove	6/26/31		
	Realty Assoc.	9/13/46		
	John White	10/14/47		
	Ridgewood Platear	10/4/49		
	Realty Assoc.	10/5/49		
	Glen Cove Realty Corp.	12/12/51		
	J. Graham	01/10/56		
	Creek Develop. Corp.	12/20/56		
	I.I. Miller	12/22/56		
	City of Glen Cove	02/13/58		
	Nassau County	04/28/70		
	City of Glen Cove	09/13/74		
	I.I. Miller	11/19/74		
•	Lee Langbaum	09/02/76		
	CONMAR Blders.	03/26/79		
	James O'Connell	12/28/79		
	Glen Cove Develop. Corp.	04/16/81		
	Village Green	08/15/83		
551 & 556	. & 556 (Similar Listings to above p			
	Glen Cove Urban Renewal	04/12/77		
	Glen Cove Community Develop.	04/16/81		
	Village Green	10/04/83		

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GARVIES POINT SITE

FIGURE 1: PROJECT SITE

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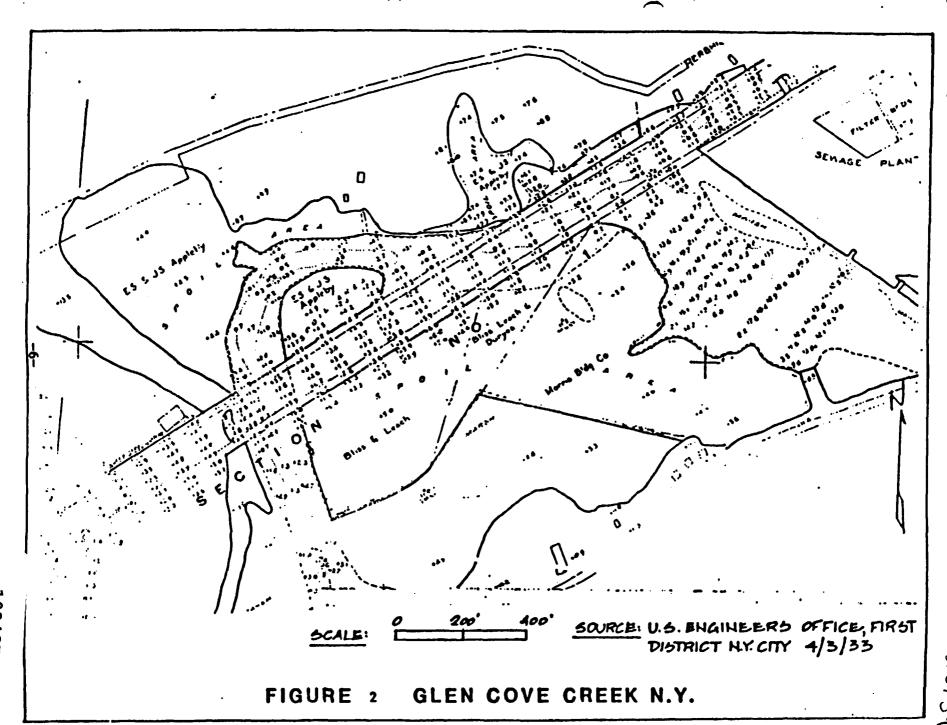
rubbish at the site and to odors allegedly coming from the uncovered sewage sludge. The City of Glen Cove was responsible for the site at this point and the City did clean and remove debris from the site on occasion.

In interviews with NCDOH staff and former staff, attempts were made to secure additional information about the site. Only hearsay information was available about possible illegal industrial dumping on the site. Although such dumping is alleged to have taken place; the types of materials, identification of what was in "drums" observed on site, and physical evidence were not carefully documented (i.e., no labels identifying the contents of drums, no surface staining was recorded and file photographs do not reveal the presence of hazardous waste). Further research into the areas of alleged waste disposal will be performed during the remedial investigation.

The Army Corps of Engineers was contacted to determine if and when dredge spoils were placed on the site. Historic records of the dredging and spoil disposal activities at Glen Cove Creek were reviewed at the Army Corps of Engineers, Navigation Section, New York District Office in New York City. Mr. S. Lew of the Navigation Section provided the files.

An April 1933 map, Figure 2, of Glen Cove Creek from the U.S. Engineers Office, First District New York City shows spoil disposal areas which implies pre-1948 dredging activity. It is obvious that the creek had a major course change prior to 1948, however, where the dredge spoils were placed cannot be determined from the Army Corps files.

Based upon the existing records, dredging of Glen Cove Creek occurred in 1948, 1960 and 1965. A review of the 1948 records determined that the volume of dredge material to be removed as 29,500 cubic yards (estimated). No maps of the actual dredging activity or spoil disposal areas were in the file.



A review of the 1960 records estimated the volume of dredge material to be removed as 27,600 cubic yards. The map of the property dated 1957, Figure 3, designates the area of spoil disposal to be in the central section of the property and calculates the area to be 7.8 acres. This designated area is within Section 21, Block A, Lot Number 471 of the City of Glen Cove. Maps of both the area to be dredged and the dredge disposal area from the U.S. Army Corps of Engineers dated 4/1/60 are shown in Figures 4 and 5.

A review of the 1965 records delineates the spoil disposal in the same area and shows the area of material to be removed to be approximately where the Glen Cove Creek discharges into Hempstead harbor.

The proposed 1979 dredging activity did not occur. The upstream area of the creek was not properly bulkheaded for dredging activity. Additionally a sediment sample of the creek obtained by the EPA showed levels of cyanide above regulatory levels. The NYSDEC subsequently declared the proposed dredge spoil as a hazardous waste and prohibited disposal of this material in the landfill. Dredging did not occur because the movement of these materials to other disposal areas was cost prohibitive.

The City of Glen Cove Building Department was contacted to determine if they possessed or had knowledge of data on disposal activities at the site. While being aware of the general nature of disposal activities on the site, the Building Department knew of no data documenting such activities.

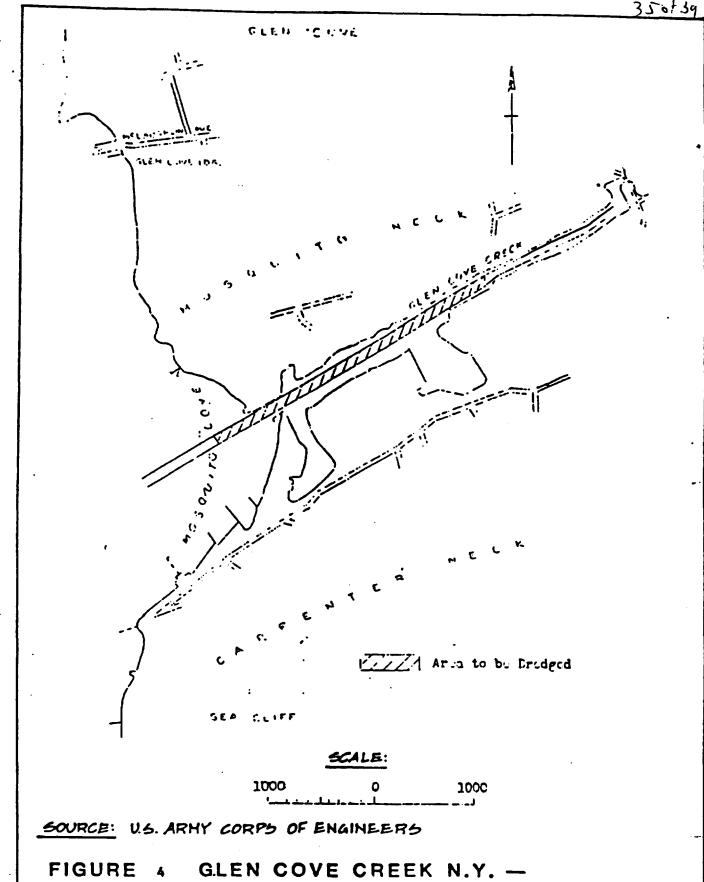
3.0 SURROUNDING PROPERTIES

The properties surrounding the site were identified during a site visit.

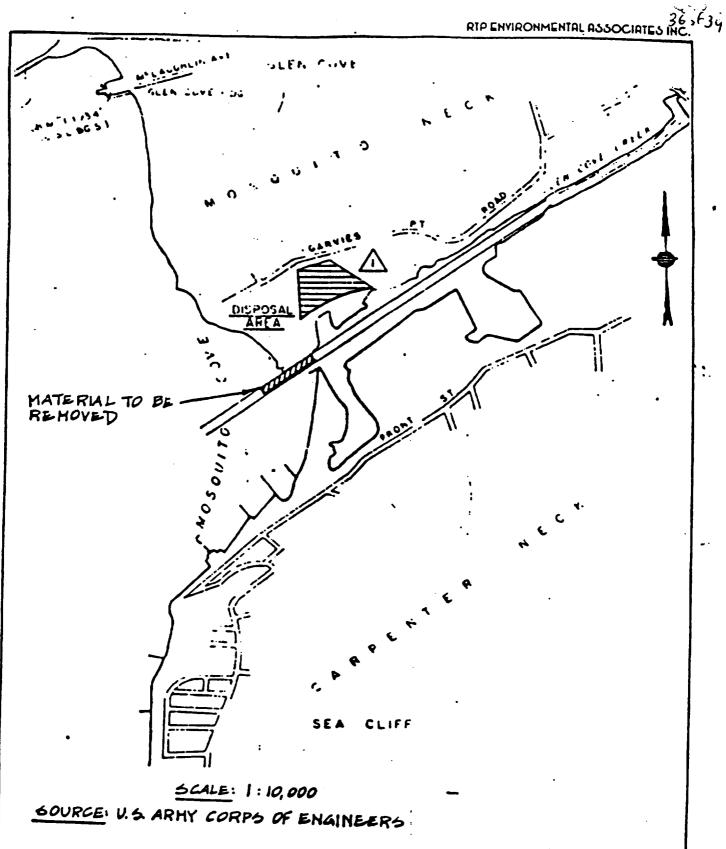
The site is bounded by Garvies Point Road to the north, Hempstead Harbor to the west, the City Anglers Club Marina to the east and Glen Cove Creek to the south (Figure 1). To the north of Garvies Point Road is the Garvies

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1960 AREA TO BE DREDGED



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FIGURE .5 GLEN COVE CREEK N.Y.-APRIL 1965 PROPOSED DREDGING ACTIVITY

Point Reserve, the Fabric Leather Corporation and the Edmos Corporation. To the west of Hempstead Harbor is the Port Washington Peninsula. To the south of Glen Cove Creek are several marinas and the City codisposal plant. To the east of the City marina is Cove Oil Company.

A search of the registered water well records at NYSDEC Region I was conducted. Wells within a one mile radius have been identified and are summarized in Table 2. The nearest water supply well on record is located at the Fabric Leather Corporation and is rated to draw 380 gallons per minute.

4.0 PREVIOUS ENVIRONMENTAL AND OTHER SITE INVESTIGATIONS

The previous site investigations are listed below in reverse chronological order. This constitutes the information available to Village Green.

Date	<u>Description</u>				
March 1986	RTP Environmental Associates and H ₂ M conduct a water test of CDM Well #2.				
Nov-1985/Feb 1986	CDM conducts a preliminary site investigation to determine if hazardous materials are present in the air, water or soils (CDM, 1986)				
1985	LKB conducts a soil boring and monitoring well program to determine subsurface soil stratigraphy, hydrogeology and soil classification. Gas monitoring wells were installed to determine subsurface methane levels.				

5.0 AERIAL PHOTOGRAPHIC INFORMATION

Three sources of aerial photographs were investigated. These included Lockwood Kessler and Bartlett (LKB), Syosset, New York; Aeroservice, Inc., Houston, Texas; and Aerographics, Inc., Bohemia, New York. Of the three

TABLE 2

PUBLIC AND PRIVATE SUPPLY WELLS WITHIN A ONE MILE RADIUS OF THE GARVIES POINT SITE

NYS DEC #	OWNER OR WELL FIELD	CAPACITY _G.P.M	DEPTH - <u>(ft)</u>	WATER
1917 2847	Wah Chang Trading Corp. Skouras Thearte Corp.	250	307	USE Processing
3466 3993	New York Water Service Corp. Henry's	425 208 20	116 177	Cooling Public Supply
4432 4440	Dykman Laundry Limco Manf. Corp.	90 200	70 352 316	General Laundry
5686 6416	City of Glen Cove Zara Asphalt Co.	45 37	92.5 106.5	General Process Water
6549-D 6587 6594	Columbia Carbon & Ribbon Zara Asphalt Co.	225 103	425 56	Diffuser
7588 7614	City of Glen Cove Hempstead Harbor Yacht Club Powers Chemco Corp.	65 -	51 26	Air Conditioning No Water
7857 8048-D	Sea Cliff Water Co. Powers Chemco, Inc.	1300	393 614 370	Diffuser Municipal Supply
8326 8327	City of Glen Cove - City Hall City of Glen Cove - City Hall	1400 1753	168 168	Diffuser Municipal Municipal
8453 8690 8709	Powers Chemco Fabric Leather Corp.	•	125 347	Test Well Cooling
8887 9612	Fabric Leather Corp. Slater Electric, Inc. Slater Electric, Inc.	380 240	312 130	Cooling
9841	Slater Electric, Inc.	240	109 121	Cooling Process Cooling

groups, LKB had the most complete set of aerial coverage of the site dating from April 5, 1950 thru March 21, 1986. Stereographic pairs are available for the following dates: 4/50, 1/55, 3/62, 566, 4/69, 4/72, 4/78, 3/84 and 3/86. The 1950 and 1955 photos are at 1"-1000' scale while the remaining years to 1984 are at 1"-1600'. The 1986 photos are at a 1"-800' scale.

Original copies of the photographs are available at Lockwood, Kessler & Bartlett, One Aerial Way, Syosset, New York 11791.

Xerox copies of the site photos for each year are attached for convenient reference.

The other sources of aerial photographs only have very limited coverages of the site. These were not investigated further because the LKB footage was considered more than sufficient to cover the period and area of interest.

6.0 OTHER SITE INFORMATION

The NYSDEC files already contain the other data that the applicant has been able to gather on the site with respect to the information request in the Order On Consent. The applicant will continue to search for additional pertinent information on the site relating to the referenced Order. These materials will also be provided to the NYSDEC according to the requirements established in the Order.

REFERENCE 4

PRELIMINARY RADIOLOGICAL ASSESSMENT
OF THE
LI TUNGSTEN FACILITY, GLEN COVE, NY

December, 1989



PRELIMINARY RADIOLOGICAL ASSESSMENT

OF THE

LI TUNGSTEN FACILITY, GLEN COVE, NY

Prepared for:

Glen Cove Development Corp. 34 Market Place Baltimore, Maryland 21202

Prepared by:

The NDL Organization, Inc. PO Box 791
Peekskill, NY 10566

December, 1989

EXECUTIVE SUMMARY

The tungsten refining process at Li Tungsten utilized ores containing low levels of natural thorium and uranium at concentrations averaging 10 to 20 pCi/g. About 2000 cubic yards of unprocessed and semi-processed ore are stored in and around the main factory on parcel A in thousands of decaying drums and crates. Ambient gamma fields near this material ranges from 10 to 30 uR/hr with levels in the 50-500 uR/hr range at 1 cm. Radon, thoron, and airborne long-lived alpha levels measured in the main (Dice) Building are within acceptable levels. The unprocessed and semi-processed ores thus do not present a significant radiological hazard.

After tungsten is removed from these ores, the radionuclides become more concentrated in the waste products which take the form of hard, concrete-like rocks, or slag, and various type of granular material. Radionuclide levels in the waste products approach 1000 pC/g and thus pose an internal radiation hazard if dusts become airborne, and an external radiation hazard by emission of beta-gamma radiation at levels in the 1 to 10 mR/hr range. There are about 500 to 1000 cubic yards of these higher level waste products piled on the ground on all 3 parcels. About 10,000 cubic yards of lower radioactivity soils and other waste materials are also piled on parcels B and C. The presence of any higher activity materials buried at greater depths under these piles is presently unknown. No significant radionuclide migration from the site via surface water run-off was detected.

A small amount of thorium metal processing also was performed at Li Tungsten. About 200 lbs of thorium metal (reading 65 mR/hr), several pails of what may be monazite or zircon sand, and 3 small furnaces with gamma levels in the 10-20 mR/hr range were found. These thorium process materials pose a greater radiological hazard than do the tungsten process materials. Fortunately, there is a small quantity of thorium materials which can be easily packaged and disposed. As of December 11, 1989, 113 cu.ft. of these highly contaminated materials and equipment have been removed from the site. Areas of thorium contamination remaining at the site include: a 1000 sq.ft. asphalted area in parcel A contaminated to 4500 dpm alpha/100 sq. cm.; a 2000 sq.ft. heavily vegetated area in parcel C with soil contaminated to about 1000 pCi/g; and a brick-lined pit behind the laboratory where thorium solutions were known to have been stored.

This survey identifies 9 specific areas of concern: 7 completely within the controlled area and 2 near the perimeter fence affecting both the controlled and uncontrolled areas. Five of these areas have been remediated in December, 1989. The data presented in this report shows pre-remediation radiation levels, however, when remediated areas are discussed, they are marked with "remediated". The 4 remaining areas include: the heavily vegetated area on parcel C discussed in the preceeding paragraph (~3-5 mR/hr), a pile of 6-12" slag rocks on the northern end of parcel C (~1-3 mR/hr), a pile of large (3 ft diameter) slag rocks along the northern fence of parcel A (~1-3 mR/hr) causing 160 uR/hr through the fence, and buried waste under the north fence of parcel A causing 300 uR/hr near the ground surface on the uncontrolled side of the fence. Remediation of these areas will require excavation with heavy equipment.

In general, the building structural components and factory equipment are not significantly contaminated. Most horizontal surfaces have an easily removable layer of ore dust. However, radiation measurements from many vats and tanks, indicate that their internal surfaces may have been contaminated with radionuclides during the various heating and chemical processing stages of tungsten refining. Closer investigation of the tank and vat interiors should be performed after their contents are characterized and removed.

Standard contamination control procedures outlined in the site safety manual will adequately protect chemical and radiological remediation teams from internal radiation exposure at the site. The remaining sources of significant external radiation (>2 mR/hr) have been roped-off.

Potential for exposure of the public is very low. The 2 areas of concern at the northern fence of the main parcel along Herb Hill Road have elevated radiation levels (0.1-0.3 mR/hr) on the grassy area between the road and the fence. Radiation levels from these sources drop off to background within 4-6 feet of the fence and pose no hazard for persons on foot or in vehicles on Herb Hill Road.

Table of Contents

EXE	CUTIV	E SI	UMM	AR!	Z		٠.				• •	• •			•		٠.	•			•			•				٠.	E-	-1
1.0	INTR	טעס	CTI	ON							• • •					• •	٠.				•			•			. •			1
2.0	OBJE	CTI	VES	• •		• •	٠.				• •				•		٠.							•						2
3.0	MEAS	URE	MEN	TS	PE	ERF	OR	ME	D		• • ·							•			•	•		•			. •		•	2
4.0	4.1 4.2 4.3 4.4	EX!	rer Pe R S	NAI TES AMI	L F STS PLE	AD S	IA	TI 	ON		U	RV:	EY	• •		• •	• •	• •	• •	• •	• •	•	 	•	• •	• •	•	• •	•	444
5.0	DISCT 5.1 5.2 5.3 5.4	EXT SUI ASS RAI 5. 5. 5.	TERI RFA SES	NAI CE SME ACT 1 E 2 C 3 S 4 C	CC ENT FIV Pro Cha Soi Con	AD NT. Ce. in l.	IAM F Y S E mi:	TI IN AI: IN Qu na	ON RB at il te	OR OR OI ib	ENE IL IL IL IL IL IL IL IL IL IL IL IL IL	E Alaispi	LS RA ND S. 	DI P	ON		CL ES	III)E M	S.AT	E	RI	AL	S	• • • • • • • • • • • • • • • • • • • •			• • • • • • • • • • • • • • • • • • • •		567778899
Figu	ire 4.		- G																В	ld	gs	3.		• 1		• •	• •	•	4 a	ì
Figu	re 4. Paro	2 - cel	Ga A:	amm Wi	na . re	Ra P	di:	at: nt	io	n Ea	Le	• V	el Bl	D dg	ia ,	ıg:	ra L	m ot	m	g	Aı	e	a.	•			•	•	4 Ł	>
Figu	re 4.	3 -	· G	amn	ıa	Ra	di	at:	io	n	Le	. V	el	D	ia	ıgı	ra	m	-	P	aı	c	el	1	3			•	4 c	;
Figu	re 4.	4 -	· Ga	amm	na	Ra	di	at:	io	n	Le	·V	e 1	D	ia	g	ra	m	-	P	aı	C	e l	(3		. •	•	4 d	l
Tabl	e 4.1		So	il	/	Pr	OC:	e 5:	5	Ma	te	r	ia	1	Ar	a	l y	si	. s	•		•	• •	•	• •			4 e	;-g	ľ
Tabl	e 4.2	2 -	Ai	rbo	rn	e i	Al	ph	a	Ra	di	.0	B.C	ti	vi	.t	Y	Me	a :	su	re	m	en	t	\$		•	• •	4	l .
	e 4.3									_		_													ĹΟ	ns	3	• •	4h	1
	NDIX																													
	NDIX											IA'	re:	RI	AI	. :	SA	MI	L	E	LC	C	ΑT	'I(NC	S				
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1.0 INTRODUCTION

The Li Tungsten facility consists of 26 acres on which exist buildings (over 300,000 sq.ft. of floor area), forested areas, ponds and swampy areas. The site was operated from the 1940's to the 1980's as a tungsten refining and processing facility and is now abandoned.

Tungsten ore, or schelite, was obtained by Li Tungsten from mines in Canada and China. In addition to tungsten, schelite contains the naturally-occurring, thorium and uranium series of radionuclides. These radionuclides are concentrated by the tungsten refining process so that the intermediate and waste products may pose a radiation hazard.

Schematics of the Thorium-232 and Uranium-238 decay series are shown in Figures 1.1 and 1.2. The Uranium-235, or Actinium, series is also present with the U-238 series, but at much lower levels. Radionuclides from these 3 series emit alpha, beta, and gamma radiation and thus pose both external and internal radiation hazards when found in high concentrations.

Approximately 12,000 cu.yards of ores and slags have been found on site. Most of this material occurs in piles on the two forested parcels of land. However, about 2000 cu.yards of material is contained in thousands of decaying drums and crates located in the main plant building.

Thorium occurs naturally in soil at a background level of about 1 pCi/g, or about 9 ppm (ref: NCRP 45). Uranium-238 and its progeny, including Radium-226, occur naturally in soil at a concentration of 0.6 pCi/g (ref: NCRP 45). Thorium concentrations in the various ore samples from the site range from background to 10's of pCi/g. Concentrations in slag, sludge, and other waste products range from 100's to 1000's of pCi/g. For comparison, the limit for unrestricted use, or "clean", soil in New York State Code Rule 38, Table 5 is 500 ppm source material, or 55 pCi/g Thorium or 180 pCi/g Uranium. The limit for radium in soil affected by uranium mines and mills is 5 pCi/g from the surface to 15 cm deep and 15 pCi/g below 15 cm (ref: EPA, 40CFR192 Uranium Mill Tailings Control Act).

Pure thorium is used in metal alloys, as optical coatings on camera lens, in vacuum tubes, and in arc lamps. It historically has been thought of as a bulk, low radioactivity material. "Source material" is a general term referring to thorium or uranium as metal, ore or bulk material. Any industrial or educational institution may possess up to 15 lbs of source material under a general license without regard to specific licensing or regulatory authority except disposal (Ref: NYS Code Rule 38 and NYS Sanitary Code Part 16). Also, possession of ores or products containing less than 500 parts per million (ppm) source material are exempt from licensing (NYS Code Rule 38).

The predecessor company to Li Tungsten, Wah Chang Trading Company, had a NYS Dept. of Labor radioactive materials license to work with pure thorium and high level thorium ores. About 10 pails and drums of material having thorium series concentrations in the 10,000 pCi/g range have been found. One drum contained about 200 lbs of thorium metal. These are likely remnants of the thorium work which should have been discontinued when the license was terminated in the early 1970's.

2.0 OBJECTIVES

The objectives of this survey are to:

- 1. assess the types, quantities, and concentrations of radionuclides present at the site;
- 2. identify radiological hazards that may be encountered by chemical and radiological remediation teams; and
- 3. identify radiological hazards that may cause exposure to the general public.
- 4. identify for subsequent removal and disposal up to 15 drums material exhibiting elevated radiation levels over 1 mR/hr.

The present project is intended to be an initial screening survey utilizing survey instruments and approximately 150 soil/process material samples.

3.0 MEASUREMENTS PERFORMED

A map of the site was overlaid with a 25'x25' grid. Gamma radiation exposure levels were measured using a NaI probe, or microR meter, at site locations within each section of the grid. Exposure rates from many radiation sources were investigated and recorded along with ambient exposure rates. A Geiger-Mueller probe in a large area configuration (pancake probe) was also used to investigate the many types of sources found. This probe is sensitive to alpha, beta, and gamma radiation and is more directional then the microR meter. The pancake probe is useful to measure specific sources in the presence of other interfering sources.

4.0 RESULTS

4.1 EXTERNAL RADIATION SURVEY

Readings from the microR and GM pancake meters are shown in Appendix A. The microR data is plotted on maps of the plant in Figures 4.1 through 4.4 as a graphical representation using cross hatching of grid sections in which elevated gamma radiation levels were detected.

4.2 WIPE TESTS

The results of wipe samples are shown in Appendix C. Only 2 of the 274 wipes showed significant removable activity.

4.3 AIR SAMPLES

Results of 4 air samples for long-lived alpha activity collected in the Dice Building are shown below:

Table 4.2 - Airborne Alpha Radioactivity Measurements

Sample#	Date	Location	Gross Alpha Conc.
RAD/A-1	6/27/89	Dice Bldg near ore pile	< 1.7x10-13 uCi/ml
RAD/A-2	6/27/89	Dice Bldg near roto kiln	< 1.8x10-13 uCi/ml
RAD/A-3	6/30/89	Dice Bldg in mixing room	< 3.8x10-13 uCi/ml
RAD/A-4	6/30/89	Dice Bldg in mixing room	< 2.8x10-13 uCi/ml
	Rule 38 li	imit	
Publ	ic:		2x10-12 uCi/ml Th-nat
Occu	pational:		6x10-11 uCi/ml Th-nat

Radon-222 and Radon-220 (Thoron) results obtained by the Dept. of Energy are shown in Table 4.3.

4.4 SOIL/PROCESS MATERIAL SAMPLING

The locations of sample collection are shown in Appendix B. The results of the samples analysis are shown in Table 4.1.

An alpha scintillation detector was also used to measure fixed alpha contamination. However, these measurements were of limited use because the source of alpha radiation, the ore, caused self-attenuation of the alpha particles, thus preventing efficient detection. Significant fixed alpha activity was found only in a few areas.

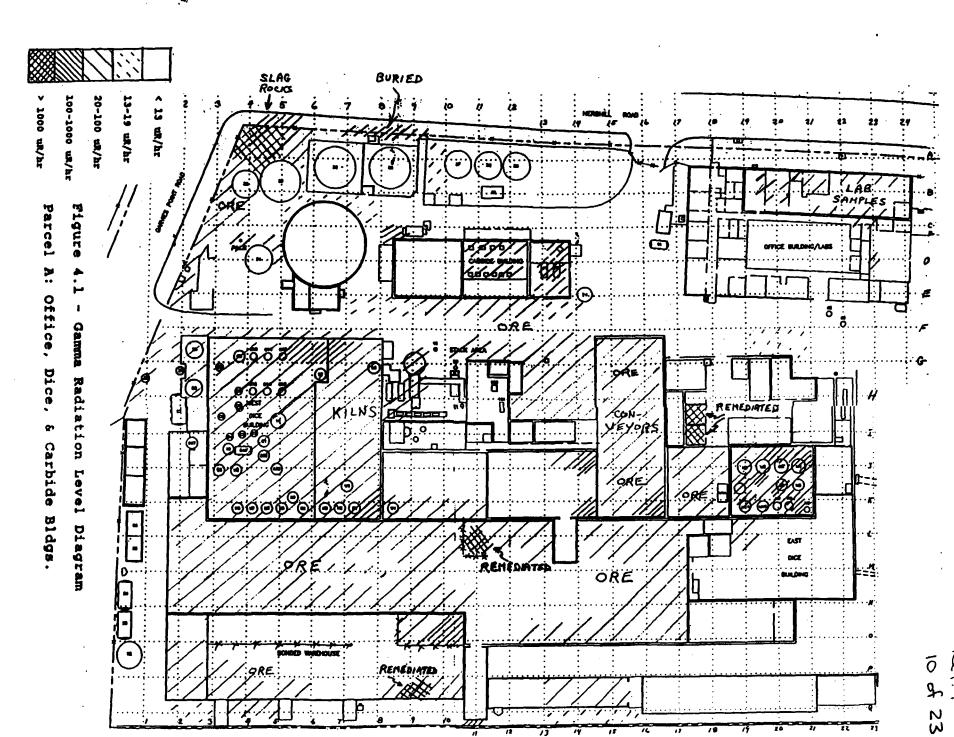
Wipe tests were performed on floors and equipment in the Offices, Laboratory, Wire Plant, and Dice Building to assess removable alpha and beta contamination. They also proved to be of limited usefulness because most of the contamination occurred in the form of low specific activity, ore dust which is not collected by wipes in sufficient quantities to be detected.

Radionuclide concentrations in solid material were investigated by the collection of 152 samples. They included samples of ores, slag, intermediate materials, pond/swamp sediment, and soil. Soil samples were collected both within the plant boundaries and around the perimeter. Normally, 3 samples were collected from each soil sampling location: surface, 6-12", and 12-24" deep. In this way, data concerning radionuclide leaching, migration or burial could be obtained.

The soil and process material samples were analyzed with a gamma spectroscopy system using a high purity germanium detector. Gamma rays from Ac-228 were used to determine the Th-232 concentration, that of Th-234 and Pa-234m were used to determine U-238, and that of Pb-214 and Bi-214 were to determine Ra-226. Gamma rays from other nuclides in the decay chains were analyzed but the results are not listed in this report unless unusual chain equilibria were found. These analyses are useful not only to compare radionuclide concentrations with regulatory guidelines, but also to gain information on the types of processes and materials that were used at Li Tungsten.

Several air samples were collected and analyzed for long lived alpha emitters. The Dept. of Energy's Environmental Measurements Laboratory also conducted thoron (radon-220) measurements in the main plant building. The purpose of air sampling and analyses was to assist in establishing appropriate levels of protection for on-site workers during implementation of remedial actions at the site.

The survey was performed during the period June to September, 1989.



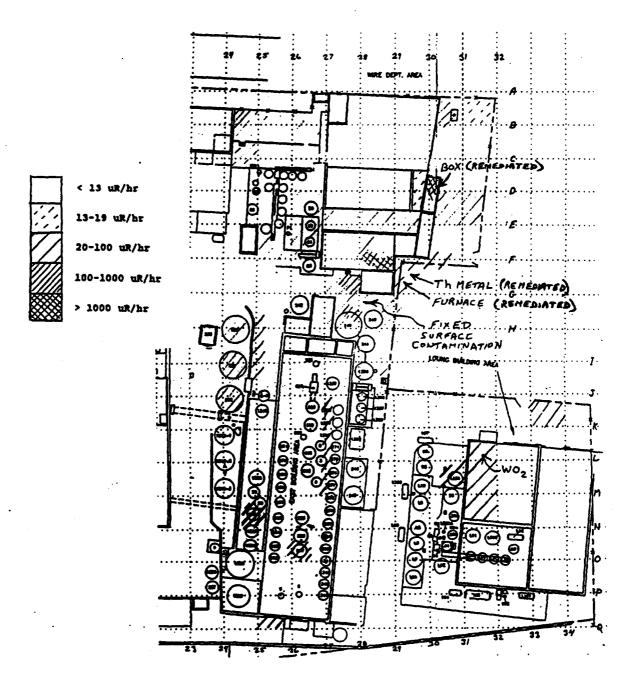
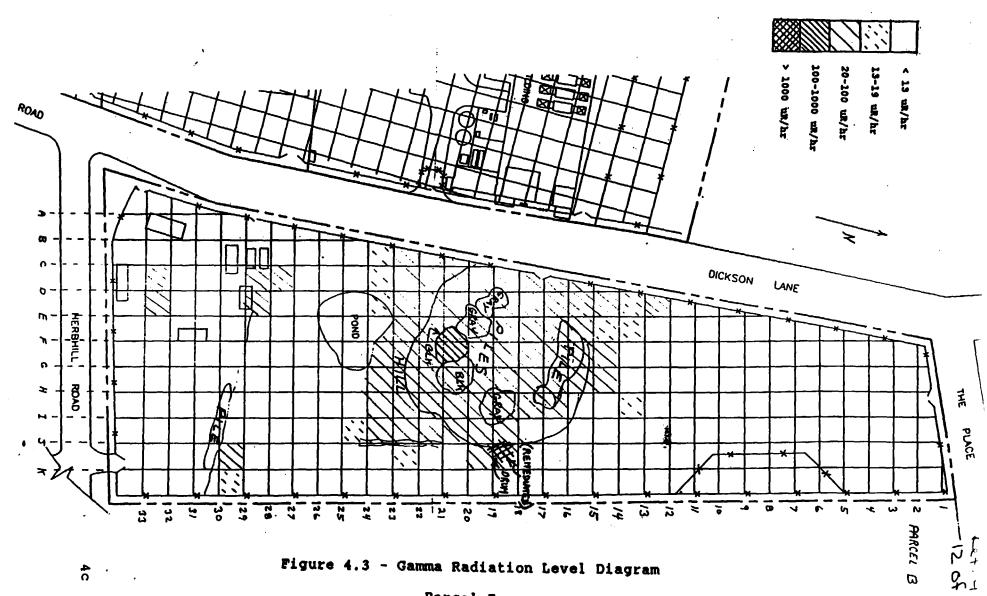


Figure 4.2 - Gamma Radiation Level Diagram

Parcel A: Wire Plant, East Bldg, & Loung Area



Parcel B

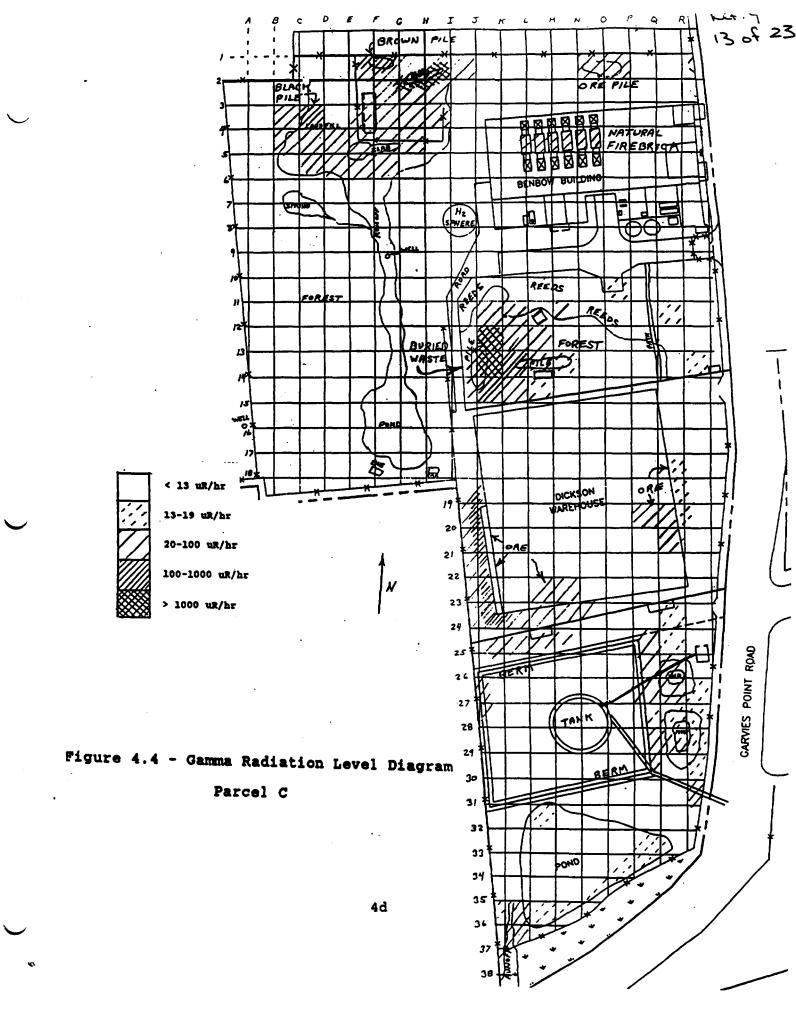


Table 4.1

LI TUNGSTEN SOIL / PROCESS MATERIAL ANALYSIS

SAIPLE 8	VT PAR- (g) CEL		TH-232+ Daughters	U-238 + Daughters	RA-226	CPH READING	uR/hr	SAMPLE DESCRIPTION
	.		(pCi/g)	(pCi/g)	(pCi/g)			
P1	105.9 8	025	1.3 ← 0.4	(1.5 ← 0.0	0.6 ← 0.2	60	10	POIND SEDIMENT IN SIDE OF POIND
P2	108.9 B	626	1.0 + 0.4	(2.1 + 0.0	0.6 +- 0.1	60	28	POND SEDIMENT - E SIDE OF POND
P3	74.7 C	L35	2.5 + 0.5	(3.0 + 0.0	1.1 + 0.2	60	10	POIND SEDIMENT - SH SIDE OF POIND NEAR OIL TANK
P4	106.0 C	N34	5.2 + 1.8	19.6 + 3.7	6.4 +- 0.5	60	14	POND SEDIMENT - E SIDE OF POND NEAR OIL TANK
P5		F17	1.6 + 0.3	(1.7 +- 0.0	1.0 ← 0.1	60	10	POND SEDIMENT - E OF DICKSON, RUNOFF FROM LANDFILL
1	145.0 A	A5	7.9 + 0.0	15.6 ← 0.0	2.5 + 0.3	200	22	BLACK ORE SPILLED - NA CORNER
2	121.0 A	B3	2.6 + 0.4	(3.3 + 0.0	2.4 + 0.2	100	22	BROWN ORE - NW CORNER
	136.5 A	B4	6.8 + 2.0	54.0 ← 5.6	64.0 +- 1.0	1000	900	BLACK ORE - NH CORNER
4	126.0 A		4.7 + 0.0	5.2 + 0.0	3.2 + 0.3	100	20	GRAY ORE - NA CORNER
5	110.0 A		9.5 - 0.0	(9.8 + 0.0)	12.3 +- 0.0	100		LT, BROWN POWDER - NEAR MEST DICE
	122.0 A		9.1 + 0.0	%.0 ← 0.0	42.1 + 0.9	600		OK. BROWN GRANDLAR - NEAR NEST DICE
	132.5 A		9.5 ← 1.6	(22.7 + 0.0)	29.9 + 0.8	600		GRAY BROWN POWDER - NEAR WEST DICE
		M	175.0 + 2.0	(90.0 + 0.0)	59.9 +- 1.1	4000	1400	large slag boulder - NJ Corner
		F9	12.8 + 0.9	8.5 2.8	8.7 ← 0.4	300	60	OK. GRAY STORE/GRANULAR
		F13	3.7 ← 0.2	(3.4 + 0.0)	3.8 + 0.1	300	20	RED/BROWN IN CRATE
	113.5 A	K17	80.4 ← 1.9	(32.0 + 0.0	(2.4 + 0.0)	1000	100	GRAY/BROWN/MITE IN CRATE PO163 DB - DICE
12	214.5 A	L16	137.0 + 2.0	(27.3 + 0.0)	(5.0 + 0.0)	1500	150	GRAY CHUNKS IN PILE - DICE
13	123.5 A	17	5.4 ÷ 1.5	(6.2 + 0.0)	6.3 + 0.4	150	20	GRAY PONDER DRUM 027 - DICE KILM BAY
14	159.5 A	K4	8.9 + 1.1	(5.6 + 0.0	6.8 + 0.4	100	40	BLACK FLOOR DIRT - NEST DICE
15	236.5 A	L3	(1.4 + 0.0)	(4.9 + 0.0)	(1.1 + 0.0)	30	20	BLACK METALLIC FINES F3 - DICE
16	288.3 A	L11	1220.0 + 5.0	<144.0 ← 0.0	52.1 + 1.7	10000	1400	GRAY SLAG ROCKS IN DRUM IN FENCE AREA - DICE
17A	191.5 A	Q11	3.9 + 0.9	(3.6 +- 0.0	1.7 - 0 .3	300	200	SOIL AT BULICHEAD DICE LOADING DOCK - SURFACE
17B	159.5 A	Q 11	3.0 + 1.1	(6.2 + 0.0)	(1.6 + 0.0)	300	300	SOIL AT BULKHEAD DICE LOADING DOCK - 6°
170	181.5 A	Q11	(4.0 + 0.0)	(4.2 + 0.0)	1.1 + 0.3	0	80	SOIL AT BULHEAD DICE LOADING DOCK - 12"
18	145.3 A	014	2.5 + 0.4	(3.6 + 0.0)	$1.7 \leftarrow 0.1$	60	8	DRAIN IN ALLEYMAY - DICE
19A	185.0 A	024	1.6 ← 0.8	(4.6 +- 0.0	1.4 + 0.2	60	6	soil at Bulkhead at East Bldg - Surface
	183.0 A	024	(2.6 + 0.0)	4.3 + 1.2	(1.6 + 0.0)	60	10	SOIL AT BULKHEAD AT EAST BLDG - 12°
19C	172.0 A	024	$1.2 \leftarrow 0.2$	$\{1.6 + 0.0$	$0.6 \leftarrow 0.1$	60	10	SOIL AT BULKHEAD AT EAST BLDG - 24°
	140.0 A	B14	5.6 + 1.1	(4.4 ← 0.0	1.8 + 0.6	60	10	SOIL NEAR FRONT GATE INSIDE FENCE - SURFACE
208	171.0 A	B 14	$1.7 \leftarrow 0.2$	(1.1 + 0.0)	0.4 - 0.1	60	10	SOIL NEAR FRONT GATE DISIDE FENCE - 12"
	162.3 A	B14	0.9 + 0.3	(1.2 + 0.0)	$0.5 \leftarrow 0.1$	60	10	soil near front gate inside fence - 24°
		AB .		30.2 +- 4.3	10.2 + 0.5	1300	400	SOIL AT HOT SPOT UNDER NORTH FENCE - SURFACE
	135.8 A		114.0 + 2.3	144.0 + 9.0	104.0 + 1.4	3000	600	SOIL AT HOT SPOT UNDER NORTH FENCE - 6°
	138.0 A			103.0 ← 8.0	103.0 + 1.4			SOIL AT HOT SPOT UNDER NORTH FENCE - 15°
22A	84.7 A		3.2 + 0.5	(3.4 + 0.0)	1.6 + 0.1	800	120	SOIL AT UTILITY POLE NEAR POOL - SURFACE
		C4	4.4 ← 0.3	(4.5 ← 0.0	1.9 - 0.1	800	120	SOIL AT UTILITY POLE HEAR POOL - 6"
	100.0 A	01	4.7 ← 1.4	(6.0 ← 0.0	(2.1 + 0.0)	60	10	SOIL AT BULICHEAD SH CORNER - SURFACE
		Q 1	3.2 ← 0.8	(3.6 ← 0.0	(2.0 + 0.0)	60	10	SOIL AT BULIDEAD SW CORNER - 6"
	215.5 A	01	(2.2 ← 0.0	(2.9 + 0.0)	(1.6 + 0.0)	60	10	SOIL AT BULIDEAD SI CORNER - 12" HIT SOLID
		61	22.2 + 1.0	10.6 ← 2.8	1.3 +- 0.6	60	50	SOIL AT WEST FENCE - SURFACE
	194.5 A	61	30.9 ÷ 1.0	(13.3 + 0.0)	(1.3 + 0.0)	80	70	SOIL AT NEST FENCE - 4" HIT CONCRETE
	163.5 A	029	2.4 + 0.2	(2.1 ← 0.0	1.3 + 0.1	60	10	SOIL AT BULINEAD LOUNGE - SURFACE
	160.0 A	029	1.8 + 0.4	(2.0 +- 0.0	0.7 + 0.2	60	20	SOIL AT BULGEAD LOUNGE - 12"
		029	8.3 + 1.4	(15.0 ← 0.0	3.5 ← 0.3	0	20	SOIL AT BULGEAD LOUNGE - 24"
	107.5 A	E31	1.6 + 0.4	(1.5 ← 0.0	0.7 + 0.1	60	15	SOIL BEHIND HIRE DEPT. SOUTH - SURFACE
	320.5 A	E31	6.9 + 0.4	(3.2 ← 0.0	1.8 ← 0.2	60	20	SOIL BEHIND WIRE CEPT. SOUTH - 12"
	162.5 A	E31	3.0 ÷ 1.0	(5.1 ← 0.0	(1.0 + 0.0)	0	40	SOIL BEHIND WIRE DEPT. SOUTH - 24"
	181.5 A	B30	2.2 + 0.7	(6.4 ← 0.0 /7.0 ← 0.0	(1.5 ← 0.0	60	10	SOIL BEHIND HIRE DEPT. HORTH - SURFACE
	137.5 A 187.0 A	B30	(2.4 ← 0.0	(7.0 ← 0.0	1.3 + 0.4	60	20	SOIL BEHIND WIRE DEPT. NORTH - 6°
4/6	10/ .V R	B30	4.1 ← 0.7	0.0 + 0.0	1.4 - 0.2	0	20	SOIL BEHIND WIRE DEPT. NORTH - 12"

28A	158.0 A	B24	3.9 + 0.4	(6.5 + 0.0)	1.7 - 0.1	60	10	SOIL BEHIND LAB HEAR SUMP - SURFACE
_ 298		B 24	$3.2 \leftarrow 0.7$	(3.4 + 0.0)	(0.9 + 0.0)	60	10	SOIL BEHIND LAB NEAR SUPP - 12"
28C		B24	1.5 ← 0.6	(3.3 + 0.0)	1.0 + 0.1	60	10	SOIL BEHIND LAB NEAR SUMP - 24"
29		B24	18.2 + 0.9	(5.8 ← 0.0	(2.6 ← 0.0	600	200	SOIL FROM BRICK SUMP BEHIND LAB
30A		627	2930.0 ÷ 10.0	(142.0 + 0.0)	(12.0 + 0.0)	100	400	YARD DIRT UNDER OUTDR STAIR AT WIRE DEPT
308		627	21.0 ← 0.0	(20.0 ← 0.0	(1.3 +- 0.0	1500	400	1' ASPHLT @ STAIR WIRE DEPT 4500 DPN/100SQCH ALPHA
31A		H27	1.7 + 0.0	(6.0 ← 0.0	2.0 + 0.2	200	50	YARD DIRT UNDER TANK BETWEEN EAST BLDG & WIRE DEPT
32A	_	H28	256.0 ÷ 2.0	(21.6 ← 0.0	(2.8 ← 0.0	200	140	DIRT/SAND AT EAST FENCE NEAR LILCO
33A		C25	3.2 + 0.7	2.7 + 1.3	0.8 + 0.2	200	20	SOIL BEHIND LAB AT END OF PORCH
34		F26	24.7 + 0.7	18.4 + 3.3	141.0 + 0.5	500	80	GRAY POWDER FROM SCOOP ELEVATOR WIRE YARD
35A		134	1.5 + 1.1	(3.3 ← 0.0	10.9 + 0.2	60	10	SOIL SE CORNER - SURFACE
358		134	2.1 + 0.5	(4.2 ← 0.0	1.2 + 0.1	60	15	SOIL SE CORNER - 6° (UNDERGRO WATER) SOIL SE CORNER - 12° (UNDERGRO WATER)
35C		K34	1.7 + 0.8	(2.2.+- 0.0	1.2 + 0.1	.60 .60	15 10	SOIL SE CURRER - 12 (UNDERSED WHICH) SOIL S SIDE - SURFACE (WET AREA)
36A 36B		F34 F34	1.7 ← 0.4 1.7 ← 0.4	(2.2 ← 0.0 (3.8 ← 0.0	1.4 +- 0.1 1.0 +- 0.1	.60	12	SOIL S SIDE - 6" (NET AREA)
36C		F34	2.5 ÷ 0.6	(2.4 + 0.0	1.7 + 0.1	60	12	SOIL S SIDE - 12" (WET AREA)
37A		A34	1.3 + 0.4	(2.7 ← 0.0	0.8 ÷ 0.1	60	10	SOIL SH CORNER - SURFACE
37B		A34	1.2 + 0.6	(2.8 + 0.0	1.0 ← 0.1	60	12	SOIL SH CORNER - 6°
37C		A34	1.3 + 0.5	(2.0 ← 0.0	0.9 + 0.1	60	12	SOIL SH CORNER - 12°
36		621		19.2 + 1.6	1.0 ← 0.1	200	50	BLACK PILE AT 2 0°C
39A		F22	3.6 ← 0.3	(6.2 ← 0.0	2.0 + 0.1	250	40	SOIL BEHIND RAD/S38 PILE - SURFACE
398		F22	2.6 + 0.4	(3.2 + 0.0	1.8 + 0.1	250	60	SOIL BEHIND RAD/S38 PILE - 6°
39C		F22	6.0 ← 0.5	(5.0 ← 0.0	2.9 + 0.2	250	60	SOIL BEHIND RAD/S38 PILE - 12°
40A		023	22.4 + 1.3	29.7 + 4.9	42.8 + 0.8	300	50	SOIL BETWEEN DROP OFF & POND - SURFACE
408		023	3.9 ← 0.7	(7.7 ← 0.0	2.6 + 0.2	250	30	SOIL BETWEEN DROP OFF & POND - 6"
40C		023	(4.5 ← 0.0	4.8 - 2.0	2.8 + 0.3	200	30	SOIL BETWEEN DROP OFF & POND - 12"
41		F21	20.1 + 0.9	187.0 + 5.0	$110.0 \leftarrow 7.0$	1000	250	BLACK PILE AT 3 O'C
42	127.7 8	E20	2.2 + 0.5	(2.9 + 0.0)	1.2 - 0.3	60	15	GRAY PILE AT 4 0°C
43	129.3 B	D20	8.7 ÷ 1.4	(12.5 + 0.0)	1.5 + 0.5	100	25	GRAY SHALE-LIKE PILE AT 5 O'C
44A	187.2 B	E18	(3.3 + 0.0)	(3.2 ← 0.0	1.1 +- 0.4	60	20	SOIL IN CENTER OF RING OF PILES - SURFACE
44B		E18	$1.7 \leftarrow 1.0$	(6.8 +- 0.0	1.3 + 0.3	60	17	SOIL IN CENTER OF RING OF PILES - 12°
44C		E18	3.5 ← 0.9	(5.2 + 0.0)		60	17	SOIL IN CENTER OF RING OF PILES - 24"
45		H19	6.0 ← 1.4	6.6 + 2.3	4.7 +- 0.4	60	25	GRAY PILE AT 12 0°C
464		[18	4.9 ← 1.2	20.6 + 3.5	20.4 + 0.6	200	40	SOIL BEHIND PILES AT 11 O'C - SURFACE
468	160.0 B		2.6 + 0.8	(8.9 ← 0.0	$3.1 \leftarrow 0.3$	100	30	SOIL BEHIND PILES AT 11 0°C - 12°
46C		I18	3.4 +- 0.8	5.5 ← 1.9	3.2 ← 0.5	100	30	SOIL BEHIND PILES AT 11 0°C - 24°
47A		F17	(1.6 ← 0.0	(2.2 + 0.0	(0.4 +- 0.0	100	18	PILE OF SOIL AT 9 0°C
48A		E15	2.2 ← 0.5	(2.0 ← 0.0	1.3 +- 0.1	60	15	SOIL BEHIND PILES AT 9 0°C - SURFACE
488			1.2 + 0.7	(1.9 ← 0.0	1.2 + 0.1	60		SOIL BEHIND PILES AT 9 0'C - 6' SOIL BEHIND PILES AT 9 0'C - 12'
48C 49A	200.0 8			(2.0 \to 0.0 25.9 \to 3.6	0.9 +- 0.1 35.3 +- 0.7	60	12 80	SOIL BERDAU PILES AT 9 0 C = 12 SOIL FROM HOT SPOT AT 11 0 C
50A	110.0 B			(3.1 + 0.0)	0.6 + 0.1	400 60	10	SOIL FROM IN CORNER - SURFACE
508	130.0 B			(1.3 + 0.0	0.7 ÷ 0.1	60	15	SOIL FROM NA CORNER - 12°
50C	157.0 8			(2.6 ← 0.0	0.7 ← 0.1	80	15	SOIL FROM NA CORNER - 24°
51A	129.3 B			(1.3 + 0.0	0.6 + 0.1	60	10	SOIL FROM HE CORNER - SURFACE
518	159.5 B			(2.2 + 0.0	0.6 + 0.1	60	15	SOIL FROM HE CORNER - 12"
510	173.6 B			(1.0 + 0.0)	0.6 + 0.1	60	15	SOIL FROM NE CORNER - 24°
52A	105.2 A			(5.5 ← 0.0	(1.5 ← 0.0	60	10	SOIL FROM HE CORNER OUTSO FENCE - SURFACE
528	205.5 A			(1.9 + 0.0)	0.4 + 0.1	60	17	SOIL FROM NE CORNER OUTSD FENCE - 12"
52C	168.0 A			(1.0 + 0.0)	0.5 ← 0.1	60	17	SOIL FROM HE CORNER OUTSO FENCE - 24"
53A		J29		(3.4 + 0.0)	(1.3 + 0.0)	60	10	SOIL BETWEEN EAST & LOUNGE BLOGS - SURFACE
538		J29		0.0 + 8.3	1.8 + 0.3	60	25	SOIL BETWEEN EAST & LOUNGE BLOGS - 12"
53C	140.5 A	J29		(9.0 + 0.0)	1.6 - 0.5	60	25	soil between east & Lounge blogs - 24"
54A	235.4 C			(5.3 + 0.0)	1.4 ← 0.2	60	10	SOIL FROM RUMOFF IN CORNER OUTSO FENCE - SURFACE
54B				0.0 + 0.0	0.7 ← 0.1	60	15	SOIL FROM RUNOFF IN CORNER OUTSD FENCE - 12"
54C				(1.1 + 0.0)	0.5 +- 0.1	60	15	SOIL FROM RUNOFF IN CORNER OUTSO FENCE - 24"
55A	254.0 C			(3.1 + 0.0)	0.6 ← 0.2	60	-	SOIL BY SE FENCE AT UNDERD OIL PIPE - SURFACE
558	283.0 C	531	(1.4 + 0.0)	(3.8 ← 0.0	(0.7 + 0.0)	60	10	SOIL BY SE FENCE AT UNDGRO OIL PIPE - 12"

55C	245.5 C \$31		(2.4 ← 0.0	(1.1 ← 0.0	60	10	SOIL BY SE FENCE AT UNDGRO OIL PIPE - 24"
56A	190.5 C S24		(3.2 ← 0.0	(1.7 + 0.0)	60	10	SOIL BY E FENCE AT GATE E OF DICKSON - SURFACE
568	174.5 C 524		$(1.5 \leftarrow 0.0$	0.6 + 0.1	60	18	SOIL BY E FENCE AT GATE E OF DICKSON - 12"
56C	188.0 C 524	(0.7 + 0.0)	(1.2 + 0.0)	$0.5 \leftarrow 0.1$	60	18	SOIL BY E FENCE AT GATE E OF DICKSON - 24"
57A	233.0 C S10	(2.1 + 0.0)	(5.1 + 0.0)	0.8 +- 0.4	60	12	SOIL BY E FENCE AT BENBON GATE - SURFACE
578	221.2 C S10	2.4 + 0.8	(3.1 + 0.0)	(1.6 + 0.0)	60	20	SOIL BY E FENCE AT BENBON GATE - 12"
57C	228.1 C 510	1.8 + 0.7	(3.2 + 0.0)	0.8 +- 0.3	60	20	SOIL BY E FENCE AT BENBON GATE - 24"
58A	163.8 C R1	(0.6 + 0.0)	(1.2 + 0.0)	0.7 + 0.1	60	10	SOIL NE CORNER - SURFACE
588	172.6 C R1	0.8 + 0.4	$\{1.1 \leftarrow 0.0$	$0.5 \leftarrow 0.1$	60	15	SOIL NE CORNER - 12"
59A	211.5 C AA2	(1.0 + 0.0)	(0.9 + 0.0)	(0.4 + 0.0)	60	10	SOIL NU CORNER OUTSO FENCE - SURFACE
598	198.7 C AA2		(1.0 + 0.0)	0.5 +- 0.1	60	15	SOIL NU CORNER OUTSD FENCE - 12"
59C	209.8 C AA2		(2.0 + 0.0)	0.5 +- 0.1	60	15	SOIL NU CORNER OUTSO FENCE - 24"
60A	212.0 C 030		(0.9 + 0.0)		60	10	SOIL IN OIL TANK BASIN - SURFACE
608	203.2 C 030		(1.0 + 0.0)	0.7 ← 0.1	60	10	SOIL IN OIL TANK BASIN (OILY) - 6°
61A	180.5 C P27	1.9 ← 0.8	(4.2 + 0.0	(1.5 +- 0.0	1000	160	SOIL FROM HOT SPOT ON OIL BASIN BERM - SURFACE
618	194.0 C P27	5.4 + 0.0	(6.4 + 0.0	1.6 +- 0.3	300	80	SOTL FROM HOT SPOT ON OTL BASTIN BERN - 12"
61C	177.5 C P27		(3.3 +- 0.0	(1.5 +- 0.0	300	80	SOIL FROM HOT SPOT ON OIL BASIN BERM - 24"
62A	95.0 C K13	948.0 ← 7.0	(113.0 ± 0.0	(23.0 +- 0.0		1500	SOIL FROM HOT SPOT IN FOREST IN OF DICKSON-SURFACE
628	179.2 C K13	278.0 + 3.0	(45.0 + 0.0	(3.8 +- 0.0	11000	3000	SOIL FROM HOT SPOT IN FOREST IN OF DICKSON - 6°
62C	124.4 C K13		(33.0 + 0.0	(3.8 +- 0.0	10000		SOIL FROM HOT SPOT IN FOREST N OF DICKSON - 12"
63A	181.0 C 013		(5.2 + 0.0	1.1 + 0.4	60	10	SOIL FROM SANDY SPOT N OF DICKSON -SURFACE
638	141.0 C 013	(3.9 + 0.0	(4.8 + 0.0	1.7 + 0.5	60	10	SOIL FROM SANDY SPOT IN OF DICKSON - 12°
63C	200.7 C 013	(3.7 + 0.0	(5.5 +- 0.0	1.1 + 0.3	60	10	SOIL FROM SANDY SPOT IN OF DICKSON - 24°
64A	247.8 C F10	1.5 ← 0.6	(2.7 +- 0.0	1.1 + 0.3	60	10	SOIL FROM LANDFILL RUNOFF NEAR SPHERE - SURFACE
64B	189.0 C F10	1.3 +- 0.0	(3.2 + 0.0	(0.7 + 0.0	60	12	SOIL FROM LANDFILL RUNDFF NEAR SPHERE - 6"
64C	207.5 C F10	(2.1 ÷ 0.0	(5.6 +- 0.0	0.9 + 0.3	60	12	SOIL FROM LANDFILL RUNGF NEAR SPHERE - 12"
65A	177.5 C C8	(2.6 + 0.0	(3.0 + 0.0	(0.8 + 0.0)	60	10	SOIL FROM SPRING S OF LANDFILL
66	207.0 C E5	2.5 + 0.7	4.6 + 1.4	3.1 + 0.3	200	20	SLAG ROCKS FROM LANDFILL S OF PROPANE TANK
67	217.5 C C5	(2.0 + 0.7	(2.7 + 0.0	1.3 + 0.3	60	12	SAND FROM LANDFILL SH OF PROPARE TANK
68	217.5 C C3	(2.2 +- 0.0	(3.8 +- 0.0	1.5 ÷ 0.2	60	10	HATERIAL FROM LANDFILL AT FENCE NEAR PROPANE TK
•••	207.0 C E3						
69		173.0 + 3.0	120.0 +- 10.0	134.0 ÷ 2.0		1000	BLACK PILE IN LANDFILL E OF PROPANE TANK BROWN PILE IN LANDFILL IN OF PROPANE TANK
70 71	200.5 C F2 161.4 C H2	38.3 ← 1.3	(18.0 ← 0.0				
71		283.0 ÷ 3.0	256.0 + 15.0	192.0 + 2.0			SLAG ROCKS FROM LONG PILE NE OF PROPANE TANK
72	235.0 C 64	(1.7 ← 0.0	(4.3 ← 0.0	1.5 + 0.2	60	20	NATERIAL FROM LANDFILL IN GULLY E OF PROPANE TANK
73A	238.5 C L2	(2.6 + 0.0	(5.4 +- 0.0	1.0 + 0.3	60	11	SOIL FRI RUNOFF BETWEEN LANDFILL & DICKSON LANE
74A	231.2 C 613	1.2 + 0.7	(2.8 ← 0.0	(1.6 ← 0.0	60	10	SOIL FROM LANDFILL RUNOFF NEAR LOWER POND - SURFAC
75A	161.0 C A18	(0.9 + 0.0)	(1.2 + 0.0	0.6 + 0.1	60	10	SOIL FROM HID-SE CORNER
76A	176.0 C A10	0.7 ← 0.4	(1.3 ← 0.0	0.5 + 0.1	60	10	SOIL FROM W SIDE AT FENCE
80A	150.0 C 027	(2.5 + 0.0)	(8.3 ← 0.0	1.6 + 0.3	100	20	SOIL FROM HUD POOL S OF WAREHOUSE
81A	202.0 C 029	6.2 + 0.8	(7.3 ← 0.0	5.1 +- 0.3	60	30	SOIL FROM 2ND NUD POOL S OF WAREHOUSE
82	185.9 C M25	11.0 ← 0.8	9.2 ÷ 3.5	10.8 + 0.4	250	20	SPILLED BLACK ORE S OF WAREHOUSE
83A	136.3 C J27	5.8 ← 1.2	(5.8 ← 0.0	7.7 - 0.4	60	20	SOIL FROM OIL TANK BERM SHI OF MARENSE, HI OF TANK
84A	178.0 C J31		(3.2 ← 0.0	0.7 + 0.3	60	10	SOIL FROM OIL TANK BERM, SW OF TANK
85A	324.7 C K37	0.9 +- 0.5	1.8 ÷ 0.0	$1.3 \leftarrow 0.1$	60	20	SOIL FROM SW CORNER POND RUNOFF INSIDE FENCE

45

TABLE 4.3

LI TUNGSTEN CO., DICE BUILDING

RADON/THORON PROGENY AIRBORNE CONCENTRATIONS

TIME	LOCATION			1-214 P		NS Bi-212	LE	RKING VEL(WL)
	·			Bq/m ³ pCi/l)			GRAB*	INTEGRATING**
8/29/89	East Dice	36.6	8.1	7.4	65.6	11.7	0.23	0.33
10:54 am		(1.0)	(0.22)	(0.2)	(1.8)	(0.32)	•	
8/29/89	West Dice	39.9	32.7	27.2	18.2	9.95	0.07	
11:39 am		(1.1)	(0.88)	(.74)	(.49)	(.27)		
8/29/89	Central Dice	39.6	32.2	20.2	14.0	3.0	0.06	0.09
12:20 pm		(1.1)	(.87)	(.55)	(.38)	(.08)	•	
8/29/89	Outside	1.72	1.3	1.1	0.01	0.03	4.0E-0	4
1:00 pm	Entrance	(.05)	(.04)	(.03)	(.0003	(80008)		

^{*} Grab filter sample -- Sampling time 30 minutes; Total WL (Rn-222 and Rn-220 progeny)

Applicable limits for non-occupational exposure:

111 Bg/m³ (3 pCi/l) Rn-222 and progeny (NYS CR 38)

370 Bq/m 3 (10 pCi/1) Rn-220 and progeny (NYS CR 38)

148 Bg/m³ (4 pCi/l) Rn-222 and progeny (EPA guidline)

The data in this table were provided by the US Dept. of Energy, Environmental Measurements Laboratory.

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^{**} Alpha Prism Integrating Monitor -- Sampling time 16.5 hours; Thoron WL (Rn-220 progeny)

5.0 DISCUSSION

5.1 EXTERNAL RADIATION LEVELS

Some materials and equipment have been found to cause gamma radiation levels of 1 to 3 mR/hr @ 1 cm and beta-gamma radiation levels estimated to be up to 65 mR/hr. Radiation levels drop off rapidly as one moves a few feet from a source. Ambient, whole body, gamma radiation fields range from 0.01 mR/hr to 0.03 mR/hr.

Eight specific locations were found to have gamma radiation levels significantly higher than other areas of the plant. These sources emitted gamma radiation levels greater than 1 mR/hr and beta-gamma levels greater than 3 mR/hr. One additional area exhibited elevated gamma levels beyond the property line. These 9 areas of concern are described below:

- 1. parcel A, grid A4: in yard area near north fence, large slag rocks ("1 meter in diameter) reading 1.4 mR/hr gamma, 10,000 cpm (approx. 3 mR/hr) beta-gamma, 175 pCi/g Th-232, and 60 pCi/g Ra-226; Gamma exposure through fence (property boundary on Herb Hill Road) is 160 uR/hr, dropping to background levels in the street.
- 2. parcel A, grid Q9 and Ll1: in Dice Bldg., 1-30 gal. drum and 6 two-gallon pails reading > 3 mR/hr gamma and 12,000 cpm (approx. 3.5 mR/hr) beta-gamma; (remediated)
- 3. parcel A, grid II7: in Dice Bldg., 1 three-gallon pail and 2 small furnaces (1'x1'x2') reading > 3 mR/hr gamma and 100,000 cpm (approx. 30 mR/hr) beta-gamma; (remediated)
- 4. parcel A, grid D30: in Wire Plant, 2 two-gallon pails and other bagged materials in a yellow radwaste box reading > 3 mR/hr gamma; (remediated)
- 5. parcel A, grid F28: in Wire Plant, a 30 gallon drum containing about 200 lbs of thorium metal chunks reading 65 mR/hr beta-gamma and 66,000 pCi/g Th-232 (ref: EPA analysis); and a furnace (3'x2'x2') reading 50,000 cpm (approx. 15 mR/hr) beta-gamma; (remediated)
- 6. parcel B, grid J19: in heavily vegetated area, corroded 55-gallon drum containing soil, reading > 3 mR/hr gamma and 40,000 cpm (approx. 12 mR/hr) beta-gamma; (remediated)
- 7. parcel C, grid H2: filled area on north portion, pile (approx 25'x 6' x 3' high) of small slag rocks reading 1.4 mR/hr gamma, 10,000 cpm (approx. 3 mR/hr) beta-gamma, 283 pCi/g Th-232, 256 pCi/g U-238, and 192 pCi/g Ra-226;

- 8. parcel C, grid J13-14: approx. 2000 sq.ft., heavily vegetated area north of Dickson Warehouse: soil contaminated with suspected pure thorium, reading 3 mR/hr gamma, 15,000 cpm (approx. 4.5 mR/hr) beta-gamma, 948 pCi/g Th-232, and no detectable U-238 or Ra-226;
- 9. Under north fence, parcel A, grid A8: contaminated soil under controlled area fence approx. 45' long x 4' wide, reading 0.7 mR/hr gamma, 300 cpm beta-gamma, 114 pCi/g Th-232, 144 pCi/g U-238, and 104 pCi/g Ra-226. Radiation levels on uncontrolled side of fence: 300 uR/hr at ground surface, dropping to background levels in the street.

5.2. SURFACE CONTAMINATION

Spilled ore and slag are not considered surface contamination because the radionuclides are trapped within the ore matrix and do not adhere to the surface on which the ore is located. Radionuclide concentrations in the ore and most slags are too low to be detected in quantities collected by wipe tests. Alpha scintillation measurements and wipe tests of surfaces covered with ore dust have not shown any significant radioactivity. The higher level slag rocks and intermediate process materials do show detectable alpha radiation emitted from their surfaces, however, the alpha levels are low relative to the beta-gamma levels emitted.

Only a few areas of the tungsten processing plants showed true surface contamination. Some steel vats in the mixing room of the Dice Building and 1 empty steel tank near the east building with an open inspection port showed total alpha contamination levels of 4000 to 8000 dpm/100 sq cm. Beta-gamma contamination of these surfaces were approximately 1.5 to 2 mR/hr at 1 cm. Contamination of these steel surfaces may have occurred from contact with the various radioactive elements while dissolved in the various acidic or basic solutions used in the tungsten refining process. Many other tanks and vats exhibiting elevated levels of gamma radiation may also prove to have internal surface contamination, however, gaining internal access to test all such items was not possible during this survey due to the presence of unknown and potentially hazardous chemicals.

More surface contamination of floors and equipment was expected in the laboratory and wire plant because about 200 lbs of thorium metal, a thorium process furnace, and thorium contaminated asphalt were found in and around the area. Only 2 wipe samples out of 274 showed any significant alpha contamination.

5.3 ASSESSMENT OF AIRBORNE RADIONUCLIDES

The four air samples collected in the Dice Building for longlived radionuclides showed no detectable activity. This is expected because there were no machinery, furnaces or other processes operating that would have caused contaminated materials to become airborne.

Radon, an inert gas, will emanate from a stationary pile of radium bearing material even when no mechanical disturbance is occurring. Thus, easily detectable radon levels were found in the Dice Building where most of the ore is stored.

The Radon-220 and 222 daughter concentrations measured by the Dept. of Energy in the Dice Building were relatively low. The highest radon-220 (thoron) daughter concentration found was 1.8 pCi/l (as compared to NYS CR 38 limit of 10 pCi/l for the general public). The highest radon-222 daughter concentration found was 1.1 pCi/l (as compared to NYS CR 38 limit of 3 pCi/l and EPA guideline of 4 pCi/l for the general public).

5.4 RADIOACTIVITY IN SOIL AND PROCESS MATERIALS

5.1.1 Process Materials

There are many different types of ores, intermediate process materials, and waste products present at Li Tungsten. Their colors, densities, and grain size vary widely as can be seen in their descriptions in Table 4.1. These materials (ores, intermediate process materials and waste products) are identified as such from visual observations of containers, labels and their proximity to various types of process equipment.

Ores and intermediate process materials (samples 1-7, 9-15, and 34) stored in and around the Dice Building average 23 pCi/g thorium-232, 27 pCi/g uranium-238, and 21 pCi/g radium-226. Typical concentrations in raw, unprocessed ore are about 10 pCi/g for all thorium and uranium chain nuclides.

Materials thought to be waste products are present on all three parcels. They occur in piles seemingly discarded in out-of-the-way locations. They vary widely in consistency from brown and black powders to hard rocks of slag ranging from a few inches to several feet in diameter. Radionuclide concentrations range from background to over 1000 pCi/g thorium in these materials. Radionuclide concentrations in the waste products are generally greater than those in raw ore because as the tungsten is removed, the mass of the remaining material decreases thus concentrating the radionuclides. Other physical and chemical processes during tungsten refining cause further concentration of certain minerals and radionuclides to very high levels (i.e., > 1000 pCi/g) in selected waste products (see next section).

About 15 to 20 large slag rocks, 3' in diameter, were found on parcel A, having a concentrations of 175 pCi/g Th-232 and 59.9 pCi/g Ra-226.

5.1.2 Chain Equilibria

Samples of unprocessed ore showed both thorium and uranium chain equilibria, i.e., daughter radionuclides were found in approximately equal concentrations to that of the parent radionuclides. Samples of intermediate process materials and waste products showed highly disturbed uranium chain equilibria and slightly disturbed thorium chain equilibria. Uranium chain disequilibrium occurs because of the dissimilar physical properties of uranium and radium, e.g., radium (a calcium analog) will float to the surface of a furnace melt with the slag. The uranium, having a density equivalent to tungsten (19.3 g/cc), will sink to the bottom. Other such separations are likely in the various chemical ore digestion processes that may have been performed at Li Tungsten. Uranium chain disequilibrium will remain for thousands of years because of the long half-lives of the radionuclides.

Thorium chain equilibrium also will be disturbed by physical and chemical processes, however, equilibrium will be restored within 60 years or less because of the relatively short half-lives of the thorium daughter nuclides. Most process material samples at Li Tungsten show thorium chain nuclides to be near equilibrium probably because they are relatively old materials (20-30 yrs).

All process materials showed detectable levels of both uranium and thorium chain nuclides. However, some soil and asphalt samples showed thorium only. This contamination has probably resulted from purification and processing of thorium metals and/or solutions not related to the tungsten process. As discussed above, after thorium purification, the daughter nuclides will grow in to equilibrium over a 60 year period. Thus, the age of a batch of pure thorium metal, or contamination resulting from such, can be estimated from the ratio of the daughters to the parent.

5.1.3 Soil

Soils tested around the perimeter of the plant contained radionuclides within normal background concentrations except for sample location 21, grid A8, parcel A. At this location, waste appears to have been buried or spilled along a 45' section of the fence next to Herb Hill Road. Because some of the contaminated soil appears to be outside of the fence, this area is of some concern. However, the contamination seems to covered by a layer of lower level radioactivity soil and sod which should prevent its spread until remediation is performed.

The soil around sample location #17 at the end of the Dice Building loading dock at the bulkhead, showed a general gamma radiation level of 300 uR/hr but near background radionuclide concentrations. The source or volume of contaminated soil may have been relatively small and thus was missed when the soil samples were collected.

One area of highly contaminated soil was found in the heavily vegetated area, north of Dickson Warehouse (sample location # 62). This area is cluttered with heavy debris such as concrete slabs, rusted drums, and piles of soil, wood, and rocks. Sample 62A (948 pCi/g thorium) consisted of rich, forest humus, not rocky slag as would be expected from observation of other hot spots. No U-238 or Ra-226 were detected in these samples. These facts indicate that a highly concentrated thorium powder or liquid were deposited there. Th-232 daughters were in about 75% equilibrium indicating that the age of the contaminant is about 30-40 years. This age range is also supported by the presence of trees growing from the contaminated area that are about a foot in diameter.

5.1.4 Contaminated Asphalt

Another location where concentrated thorium was thought to have been spilled is on an asphalted area between the wire department and the East Building (Sample locations 30, 31 and 32). Fixed alpha levels on the asphalt were about 4500 dpm/100 sq cm and gamma levels about 400 uR/hr at 1 cm. No U-238 or Ra-226 were detected in the samples. The surface of the asphalt was mostly free of ore dust.

5.1.5 Potential for Radionuclide Migration

Three pond, or swampy, areas exist on the site from which five sediment samples were collected. Samples Pl (parcel B, grid D24) and P2 (parcel B, grid G26) did not exhibit elevated levels of radionuclides. The presence of contaminated soil and waste piles uphill and adjacent to the pond suggests that there is very limited contaminant migration away from the piles. A similar situation exists at the filled area on the north portion of parcel C. While a large quantity of contaminated materials exist there, samples collected from run-off swales (sample 73, grid L2; sample 72, grid H4; sample 64, grid G10) and a downhill pond (sample P5, parcel C, grid F17) showed background levels of radionuclides.

Sediment samples P3 (grid L35) and P4 (grid P34,35) from the pond near the large oil tank on parcel C did should slightly elevated concentrations. Soil samples 84 (parcel C, grid K37) and 54 (parcel C, grid K38) from the run-off swale southwest of the pond, indicate that radionuclides were not being significantly transported from the site via surface water run-off.

5.1.6 Materials in Large Tanks

Several large tanks (5000 - 20,000 gallon cap.) emit gamma radiation levels of 30 to 100 uR/hr. They include tank numbers: 231, 232, 233, 1334 (empty), L138, 287, K4, K5, K6, K7, K8, and K9. Some are constructed of wood, others of steel. These readings indicate that contaminated liquid, contaminated sludge, or contaminated interior walls are present within the tank. Other tanks in the East Dice and West Dice Buildings may also emit elevated levels of gamma radiation, however, higher levels of gamma radiation from nearby piles of solid process materials may have masked radiation emitted from the tanks.

Samples of the liquid contents of many tanks were collected by Direct Environmental, Inc, in November, 1989. Gamma spectroscopic analyses show very low radionuclide concentrations (see Appendix D). The three outdoor, wooden tanks (231, 232 and 233 on parcel A, grid 24-I,J,K), reading 80-100 uR/hr, are filled with what appears to be clean rainwater. Thus, the walls themselves must be the source of the radiation. Tank # 1334 (parcel A, grid 24N), an empty steel tank with an open port, had slightly contaminated internal walls reading about 1000 cpm on the GM pancake probe and 1600 dpm/100 sq.cm. total alpha. Sludge samples from the tanks have not be studied at this time.

REFERENCE 5

RECORD OF TELEPHONE CONVERSATION

	DATE	5/3/95
TO Captain's Cove Condominium Site		10:45
NAME/FILE NO.		
FROM Mike Hustran		·
CLIENT/PROJECT E PA ARCS II		
SUBJECT Wastes from Li Tungsten disposed at Captains Go	•	
CHARGE: DEPT. NO OFS N		
DISCUSSION WITH Frank Peny - Former Litangsten (5/6) 759 - Employee	9896	

- -Mr. Pena wo. Ked for Li Tungsten from 1974 until the operations closed in 1985.
- He worked in the screening department and curbide building.
- He personally never saw wask being disposed at the Captains Gove condo site, on Garvies Point as he referred to.t.
 - During his employment at Litingsten he spoke with many of the note times" who worked at the site for many years. Many of the employee's that worked at the site in the 1950s and 1960s stred they routinely dumped wasteslay etc. at the barries point site.
 - He stated that exployers used to take boxes or crates of material (slay the), using a fack lift, drive down the road and dump the material into the landfill.

RECORD OF TELEPHONE CONVERSATION

			DATE .	5/3/95	
TO <u>Captain's Cove</u> FROM M.K. Heffre	Comminum 5	; Le		10:45 2 of Tolecon	-
CLIENT/PROJECT	•	•			
SUBJECT			· · · · · · · · · · · · · · · · · · ·		
CHARGE: DEPT. NO	CLIENT SYMBO	x	OFS NO		
DISCUSSION WITH					=
- Wastes used to be across the street and hydrogen	e dumped in 1h t by The wood tak.	ls, backby	the reduction	building,	
- He stated that wh	rile he worked	at Lilung	sten, nathral	was	
dumped on The 9	round or into 6	Hen Gove Cre	eK.	1 01/1	
dumped on The go . Large crock with whom dumped r	k or open vats ofthere or aceton	used for d	egreasing parts	to was tilled,	
waste we	TO GOTO O THE	<i>p.y.</i>		•	
(neek.			· T /	and almost is of	بمريضا
- Mr. Pena stated the with the workers /	int the owners/of	heators of L	players du	mping up to	
Creek. - Mr. Pena stated the workers! 50,000 gallons of the owners since	it cost seven	thousand doll	as for The s	sida hydroxido	, ?
Plakes.	1 /	Mr. Com	11 of EPA	to walk over	The
The owners since Plakes. -Mr. Pena stated that Li Tungsten site and Mr. coholly (212) 637	discuss the form -4291	ler disposal	practices.		
<i>1</i> ~		•			
CC:	BY	NAME	π	TLE DEPT. NO	5 .

REFERENCE 6

02-9003-01-SI REV. NO. 0

FINAL DRAFT
SITE INSPECTION REPORT
LI TUNGSTEN
GLEN COVE, NEW YORK
VOLUME 1 OF 5

PREPARED UNDER

TECHNICAL DIRECTIVE DOCUMENT NO. 02-9003-01 CONTRACT NO. 68-01-7346

FOR THE

ENVIRONMENTAL SERVICES DIVISION
U.S. ENVIRONMENTAL PROTECTION AGENCY

SEPTEMBER 28, 1990

NUS CORPORATION SUPERFUND DIVISION

SUBMITTED BY:

RICHARD FEINBERG PROJECT MANAGER

STEVEN OKULEVICZ

SITE MANAGER

REVIEWED/APPROVED BY:

RÖNALD M. NAMAN FIT OFFICE MANAGER

02-9003-01- SI Rev. No. 0 कि. ५ ३,६५३

SITE INSPECTION REPORT: LEVEL III

PART I: SITE INFORMATION

1.	Site Name/Alias			ng Smelting and Refinir	
	Street 63 Herb		n Chang Teledyn	e Inc./National Recondit	tioning Company
	City Glen Cove		· · · · · · · · · · · · · · · · · · ·	State New York	Zip <u>11542</u>
2.	County Nassau			County Code 059	Cong. Dist. 3
3	EPA ID No. NYD	986882660			
4.	Block No. 21A a	and 31G		Lot Nos. 21-A-14, 15, 495, 544, 545	
5.	Latitude 40° 51	′36″ N		Longitude <u>73° 38′ 25″</u>	<u>w</u>
	USGS Quad. Sea	Cliff, New York			
6.	Owner_Glen Co	ve Development (Company	TelNo. <u>Unavailable</u>	
	Street 34 Marke	et Street			
	City_Baltimore	5-31 -		State_Maryland	Zip <u>21202</u>
7.	Operator Li Tuno	sten		Tel. No. <u>(516) 676-1313</u>	
	Street 63 Herb H	Hill Road			
	City_Glen Cove			State_New York	Zip_11542
8.	Type of Ownersh	nip			•
		☐ Federal	☐ State		
	□ County	☐ Municipal	Unkno	own 🗆 Oth	ner <u>NA</u>
9.	Owner/Operator	Notification on F	ile		
	☐ RCRA 3001	Date NA		CERCLA 103c Da	ate <u>NA</u>
	None	☐ Unkno	own		
10.	Permit Informati	on			
	Permit	Permit No.	Date Issued	Expiration Date	Comments
	SPDES	NYD008249	Unknown	1987	Cooling Water <u>Discharge</u>
	Radiation Source Material License		<u>3/19/64</u>	Cancelled 1971	License to store, transport, and deliver radioactive compounds

UZ-9003-01- 31 Rev. No. 0 Air Disch

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narges elting	3.548

	Air Permit	Unknown	Unknown	Unknown	operation
11.	Site Status				
	☐ Active	⊠Inactive		Unknown	
12.	Years of Operat	tion <u>1941</u>	to _June	1985	
13.	above- or belov	pes of waste sourd w-ground tanks or bers as needed to i	containers, land	treatment, etc.) on s	nt, piles, stained soil site. Initiate as man

Waste Sources

Waste Unit No.	Waste Unit Type	Facility Name for Unit
1	Drums	55- and 30-Gallon Drums
2	Piles	Waste Piles /Mounds
3	Crates	Wooden Crates
4	Tanks	Tanks
5	Surface Impoundments	Mud Pond/Mud Holes/Oil Recovery
		<u>Sumps</u>
6	Landfill	Landfill
7	Stained Soil	Stained Soil
8	Buried Surface Impoundment	500,000-Gallon Fuel Oil Tan's

(b) Other Areas of Concern

Identify any miscellaneous spills, dumping, etc. on site; describe the materials and identify their locations on site.

There are five other areas or items of concern at this site. First, there is a radiation hazard. The facility smelted monazite sand and tungsten ore (scheelite/wolframite), which contain naturally radioactive thorium-, uranium-, and radium-bearing compounds, to produce tungsten carbide powder and other tungsten-containing products. In addition, commercially prepared thorium oxide, thorium nitrate, and uranium (uranyl) acetate were used during ore processing. These radioactive compounds are present in the crates, piles, drums, and landfill areas on the site in various concentrations. A previous radiation survey of the facility <u>conducted by Enviropact Services in 1988 determined gross alpha radiation of three samples of</u> unknown media from various waste containers to range from 64 to 251 nanocuries per gram (nCi/q). Another survey, conducted by the NDL Organization in 1989 for the entire site. detected various levels of radiation, with the highest level detected at 1,000 picocuries per gram (pCi/q) in tungsten waste products. Background radiation levels in soil for New York State are 55 pCi/g for thorium and 180 pCi/g for uranium. Some of the large process solution vats and equipment in the facility are also radioactive. Soil by the fence along the southern boundary of Herb Hill Road is also radioactive, with levels of 160 microRoentgen per hour (uR/hr) to 300 uR/hr from material either on the other side of the fence or buried below the fence (Ref. Nos. 2, 3, 13, 29, 33, 37).

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The second area of concern is the Dice and East Buildings. Both buildings contain many crates and stacks of 55-gallon drums and wooden barrels of raw and reprocessed ore material. Rainwater has flooded both of these buildings to a depth of approximately 1 foot; this water may also contain dissolved heavy metals and be slightly radioactive from contact with the ore material (Ref. Nos. 4, 13, 31, 34, 35, 37, 49, 51, 52).

The third item of concern is asbestos. This material is found in siding shingles, roofing tiles, tank covers, and pipe wrapping. All of these items are in a state of decay and pieces of asbestos-containing materials have been found on the ground (Ref. Nos. 4, 13, 25, 26, 48, 52).

The fourth item of concern is the empty 55- and 30-gallon drums. Many of these drums are found scattered in disordered piles and stacks throughout the site; some of these drums, though empty, may be radioactive and create both a chemical and physical hazard on the site (Ref. No. 25).

The fifth and last area of concern is the Glen Cove Landfill, located on the south side of Garvies Point Road near the Li Tungsten facility. According to the City Historian for Glen Cove, this area (Section 21, Block 259, Lot 1) served as a municipal landfill and may have received waste ore and other waste materials from the Li Tungsten facility. Analyses of soil samples collected from this area by the Nassau County Department of Health revealed above background levels of radiation. The construction of a condominium project was halted due to the discovery of radiation and hazardous waste in the area. This area should be inspected and sampled for radioactive and other hazardous waste (Ref. Nos. 30, 36).

14. Information available from

Contact Amy Brochu Agency U.S. EPA Tel. No. (201) 906-6802

Preparer Steven Okulewicz Agency NUS Corp. Region 2 FIT Date Sept. 28, 1990

Der. 6

PART II: WASTE SOURCE INFORMATION

Drums

There are 3,850 55-gallon drums and 4,333 30-gallon drums on site that contain solid, sludge, and liquid materials; some drums contain raw and processed tungsten ores and residues. The total number of drums containing solid waste is 8,052; another 131 drums contain liquids. The total capacity of these drums is approximately 341,740 gallons. The majority of the drums on site are known to contain radioactive ores and residues including uranium, thorium, radium, organics such as carbon tetrachloride, perchloroethylene, and PCBs, and inorganic materials which include lead, tungsten, chromium, cadmium, arsenic, copper, nickel, zinc, barium, hydrochloric acid, hydrofluoric acid, nitric acid, and cyanide. The drums are scattered around the site and some are clustered in several buildings. Some are suspected to be buried within the landfill area, while others are stacked within or around the Dice Building, the Dickson Warehouse, the north and south sides of the Carbide Building, and at the southern corner of Herb Hill Road and Garvies Point Road. Some of these drums are overstacked, some have toppled, some are badly corroded, and some are leaking their contents upon the ground either within or around many of the buildings on site. The condition of the drums suspected to buried within the landfill area is unknown (Ref. Nos. 2, 4, 22, 26, 49, 50, 52). Figures 1 and 2 provide a site location map and a site map, respectively. Figure 3 provides a monitoring well location map.

Waste Piles

There are nine waste piles located on the site. Seven black and grey waste piles are located around the natural pond in the landfill area between Herb Hill Road and The Place. One mound of waste is located behind and to the west of the Reduction Building and another waste pile is located north of the Dickson Warehouse. All of these piles are uncovered and there is no containment. The total volume of these piles is estimated to be greater than 325 yds³; the quantity of hazardous waste within these piles is unknown. The physical states of the waste within these piles are solids, powders, and sludges. The specific substances known to be present in these piles are the ores and residues of tungsten processing. These substances include lead, chromium, barium, copper, zinc, arsenic, cadmium, nickel, uranium, radium, thorium, and cyanide. All of the piles are known to contain radioactive compounds of uranium, thorium, and radium. The piles adjacent to the Dickson Warehouse and to the west of the Reduction Building have been roped off and marked with placards that indicate hazardous radiation; the seven other waste piles are unmarked and are not roped off (Ref. Nos. 2, 4, 22, 29, 37, 49, 52).

Crates

There are 719 wooden crates on site whose volume is estimated to be 705 cubic yards. These wooden crates are found in various areas of the site, but are located mainly within the Dickson Warehouse Building, on the north side of the Carbide Building, and within the Dice/ Warehouse Building. Some

Rev. No. Per. 6 \boldsymbol{D} Mott Point (QUAD) SEA CLIFF, N.Y.

SITE LOCATION MAP

LI TUNGSTEN, GLEN COVE, N.Y.

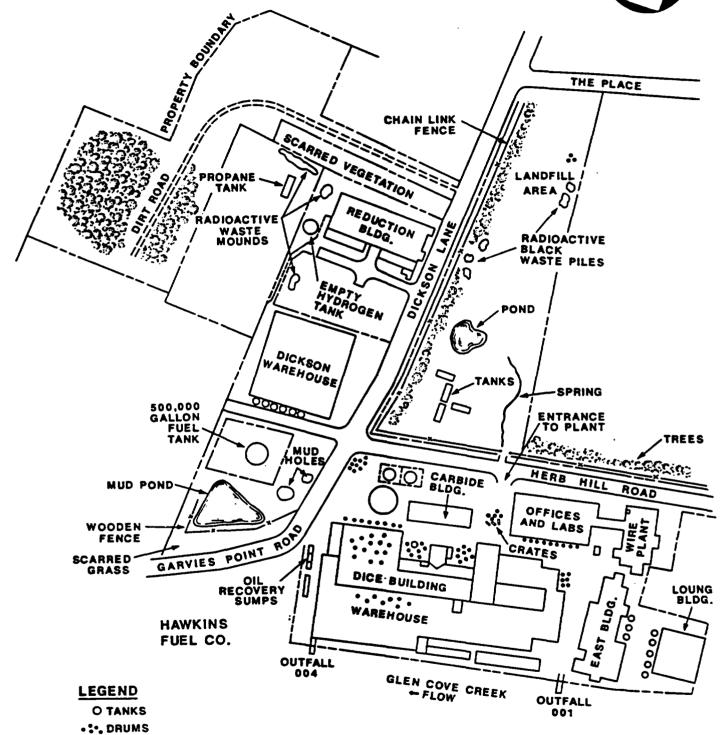
SCALE: 1'- 2000'

FIGURE 1



Rev. No. 0 Rev. 6



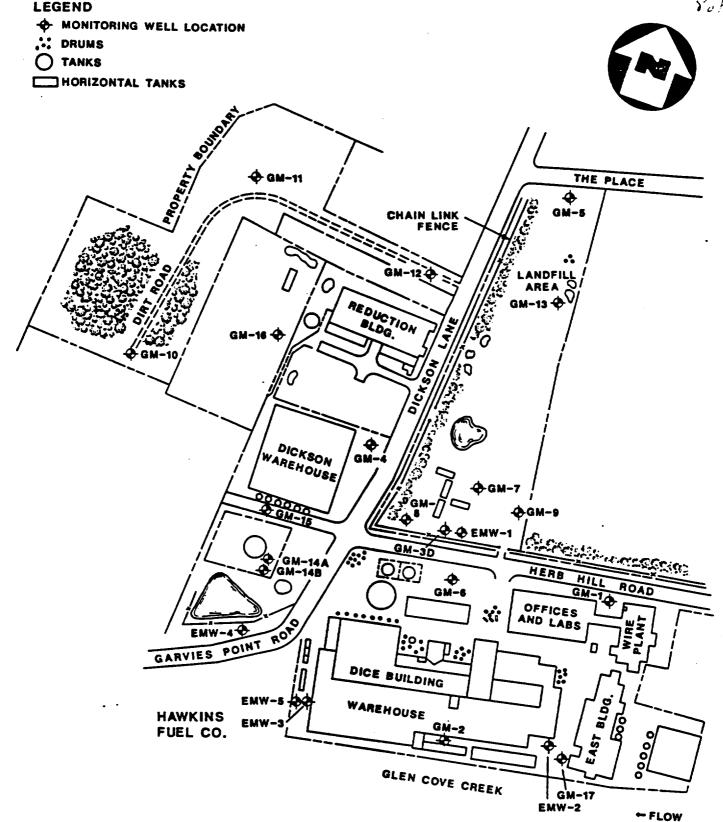


SITE MAP
LI TUNGSTEN, GLEN COVE, LONG ISLAND, N.Y.

NOT TO SCALE

FIGURE 2





MONITORING WELL LOCATION MAP
LI TUNGSTEN, GLEN COVE, LONG ISLAND, N.Y.

NOT TO SCALE

FIGURE 3



of these crates are located in open, uncovered areas outside of the buildings; these crates have been observed to be badly weathered or collapsed and spilling their contents upon the ground. Specific hazardous substances known to be present in these wooden crates are raw and processed tungsten ores that contain heavy metals including uranium, thorium, radium, lead, cadmium, chromium, copper, arsenic, zinc, nickel, and barium (Ref. Nos. 4, 13, 26, 37, 49, 52).

Tanks

There are 224 tanks made of wood, metal, or fiberglass on site. The majority of these tanks are located in the Dice Building, the Warehouse Building, the East Building, the Loung Building, to the west of the Dice Building, at the southern end of the landfill area, and to the northwest of the Carbide Building. A large aboveground 500,000-gallon fuel oil tank is located to the north of the Mud Pond. There are also two 275-gallon fuel oil tanks and one 200-gallon fuel oil tank present on site. The total capacities of 86 other tanks found to have contained liquids was estimated to be 518,131 gallons. The volume of liquids actually present in these tanks is unknown; the volume contained in 51 tanks from which samples were collected was estimated at 373,000 gallons. Two pressurized tanks also remain on site; one contains aqueous ammonia and the other contains propane gas. The volume of gas remaining in these tanks is unknown. The remaining 132 tanks either contain residual solids or are empty. The physical condition of some of these tanks is unknown. Many of the tanks are corroded or have collapsed linings. Fifty tanks have been inspected internally and externally for leaks or rupture. The contents of two tanks determined not to be secure have been sampled, drained, and drummed for disposal by Hart Environmental Consultants. None of these tanks are diked or have any secondary containment structures. The specific hazardous substances known to be present within these tanks include ammonium paratungstate (APT), ammonium hydroxide, spent hydrochloric acid, hydrochloric acid, aqueous ammonia, sodium hydroxide, tungsten acid, calcium chloride, cobalt chloride solution, sodium tungstate solution, and process solutions containing heavy metals that include arsenic, chromium, lead, thorium, tungsten, and radium. There are also approximately eight underground tanks at unspecified locations and of unknown integrity on site (Ref. Nos. 4, 13, 25, 26, 41, 49, 52).

Surface Impoundments

There are six surface impoundments on the site: two unlined settling ponds, referred to as the Mud Holes, a lined settling pond known as the Mud Pond, and three concrete oil recovery sumps. The former three impoundments are located immediately south and southeast of the 500,000-gallon fuel oil tank along Garvies Point Road. The exact volumes of the Mud Pond and Mud Holes are unknown; the quantity of waste in them is also unknown. The Mud Pond was lined with a plastic/rubber liner, but has been leaking into the groundwater and surface soil, causing scarred vegetation. A plume of waste/process water which contains heavy metals has been detected in the vicinity of the Mud Pond and the Mud Holes.

The three concrete oil recovery sumps are located west of the Dice/Warehouse Building and are connected via pipes to the Mud Pond/Mud Holes. None of these impoundments are covered. The total area of these impoundments is estimated to be 11,760 ft². The hazardous substances known to be present include sludges, fines, slurries, and liquids that contain lead, chromium, cadmium, arsenic, beryllium, antimony, cobalt, copper, manganese, nickel, zinc, sulfate compounds, chloride compounds, and PCBs (Ref. Nos. 4, 21, 22, 26, 46, 49, 50, 52).

Buried Surface Impoundment

A buried surface impoundment was located in the vicinity of and under the present location of the 500,000-gallon fuel oil tank on Garvies Point Road. The only documentation of this impoundment is in an aerial photograph of the site from 1950 (Ref. No. 46). This impoundment is suspected to have received waste/process waters from the Li Tungsten facility prior to the construction of the Mud Pond, Mud Holes, and fuel oil tank. The area of this impoundment, as measured from the aerial photograph, is approximately 5,100 ft². This impoundment is suspected to have contained contaminants similar to those presently found in the Mud Pond/ Mud Holes.

Landfill

The unlined landfill is located in an open, uncovered, partially wooded lot between the north side of Herb Hill Road and The Place. Most of the landfill is located to the northern end of the lot closest to The Place. The estimated volume of the landfill area is approximately 6,000 yds³. The actual quantity of hazardous waste within the landfill is unknown. The specific hazardous substances suspected to present in the landfill are the residues from the tungsten ore extraction process. Buried drums of unknown structural integrity were reported to be present in the area (Ref. Nos. 4, 36, 49).

Stained Soil

Stained soil is found along the perimeter of the Mud Pond/ Mud Holes and extends under and beyond the wooden fence to the edge of Garvies Point Road. Stained soil is also found around the nine radioactive waste piles. The quantity of hazardous substances present within the stained soil is unknown. The volume of material to be removed from the Mud Pond/ Mud Holes area is estimated to be greater than 5,000 yds³. The substances known to be present in the stained soil around the Mud Pond/Mud Holes include chloride compounds, sulfate compounds, No. 2 fuel oil, and heavy metals such as lead, chromium, cadmium, arsenic, and tungsten. The stained soil in the aforementioned area also has an organic odor.

Ref. Nos. 2, 4, 5, 8, 21, 26, 30, 31, 32, 37, 40, 41, 42, 49, 50, 52

PART III: SITE INSPECTION SAMPLE RESULTS

NUS Corporation Region 2 FIT conducted a site inspection at the Li Tungsten facility on April 18-19, 1990 and on May 15, 1990, during which a total of 9 groundwater, 13 surface water, 9 sediment, and 11 soil samples were collected. These samples were collected to determine whether any CERCLAeligible compounds are present in the groundwater, surface water, sediment, or soil that can be attributed to the waste units present on the site. All sample locations are shown on Figure 4 of this report. These samples were analyzed under the Contract Laboratory Program (CLP) for Target Compound List (TCL) organic and inorganic compounds, including cyanide. Refer to Table 1 in Part III of this report for a summary of the significant organic and inorganic compounds that were detected on the site. All CLP analytical data sheets are provided in Reference No. 50 of this report. Eleven surface water, 8 sediment, and 10 soil samples that had been collected by NUS Corporation Region 2 FIT in April and May 1990 were received by the National Enforcement Investigations Center (NEIC) on June 18, 1990. These samples were analyzed quantitatively for tungsten and qualititatively for copper, zinc, arsenic, molybdenum, antimony, lead, bismuth, thorium, and uranium, using inductively coupled plasma mass spectrometry (ICP/MS). Analysis for tungsten is not part of the routine analytical services performed under the CLP. Refer to Reference No. 31 for a summary of the NEIC analytical results.

02-9003-01-SI Rev. No. R.F. 6

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SURFACE WATER SAMPLE SURFACE WATER SAMPLE . DRUMS SEDIMENT SAMPLE SGROUNDWATER SAMPLE O TANKS THE PLACE GW2 S10 200 REDUCTION **S**5 PROPANE PLOG. TANK SOICK SON \$0 **SWB** SED9 SW9 DICKSON WAREHOUSE SED8 **S8** GW3 4 SED2 00 SW1,SW13 **SW4** SED4 OFFICES AND LABS S1. S11 = POINT

BUILDING

GLEN COVE CREEK

WAREHOUSE

SW10

OUTFALL 004

B SOIL SAMPLE

LEGEND

A SEDIMENT SAMPLE

SAMPLE LOCATION MAP

* SW1/3

GW8 +

SED6 SW6

GARVIES

HAWKINS

FUEL CO.

* SAMPLES COLLECTED 5/15/90

GW9 **GW10** * SED1/2

LI TUNGSTEN, GLEN COVE, LONG ISLAND, N.Y.

NOT TO SCALE



400

0000,

* SW2

GW7

Š SED7

SW7

OUTFALL 001

estimated value, concound present below CPOL but above IDL "Analysis did not pass EFA 0A/OC Presumptive evidence of the overence

- analysis not required
- tracts ale stat of fullifier

of the esternal

ILATTLES	,														
apple ID No. affic Report No.	NA3F-CMF	NYJE-GW4 BEB21	NYJL-645 06022	Ni JL-GN7 BE626	86827	BEB28	Najl-Suzins/HSD) Dedza	06030	06031	0(833	BE840	86842	0(843	86844	06046
lri:	WATER	WATER	WATER	WATER	WATER	WATER	RJIAN	WATER	WATER	WATER	WATER	SEDIMENT	SEDIMENT	SEDEMENT	SEDIMENT
its Iution fector	wg/l 	ug/L 2.78	ug/L S	ugil l	∪3/L 1	ug/l i	₩9/L l	ugit I	ug/L I	ug/L L	⊎g/l 1	ug/+g 1	ug/kg l	ug/kg 1	ug/kg l
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ns-1.2-Dichloroethene (total)	:	87								J					
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D.T.C

MPLING PATES: 64/18/90-04/19/90 'A CASE NO.: 15907 LAB: COMPUCHEN

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

ATILES ple ID Mo. ffic Recert Mc. ri. ts ution Factor cent Moisture	NYJL-SI BEBSO SOIL Ug/kg I 27	NYJL-S2(HS/HSD) OCOS1 SOIL Ug/kg 1 52	NYJL-53 BEB52 SOIL wg/kg I 19	MYJL-S4 BEBSS SOIL Ug/Fg I I:	#YJL-59 86858 501L ug/+3 L 1	NYJL-S11(DUP) BEB60 S01L Ug/) g 1 24	NYJL-RINE OCOGI WATER Ug/L I	MrJL-N1H2 BEO62 MATER Wg/L f	NYJL-RINS OEB6S HATER Ug/L I	Nyje-Rina OE064 WATER Ug/l I	NYJE-TRBI BEB69 WATER Ug/L I
Ordaelhane	•••••••	****************	••••••		• • • • • • • • • • • • • • • • • • • •		•••••	••••••	•••••		
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·Dichloroethane	•	••									
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2-Trichloroethane											
ene	•										
ns-1.3-bichloroprocene	•										
ofore	į	•									
thy1-2-Fentanone	:										
#anone	:										
achlorcethene	i										
ene	į	•									
2.2-letrachloroethane	i										
robenzene	i										
lbenzene	i										
ene	i										
ne: llotal)	•										

ik space compound analyzed for but not delected compound found in lab tlank as well as sample, indicates possible/trobable blank contamination estimated value estimated calue, compound present below CROL but above IDL analysis did not pass EFA 9A/9C Fresumetive evidence of the presence of the material analisis not required ction limits elevated if fillution

). 100			. 11	Mhiu	ne	J. In) 113445	Lin ANA	drucA	L VAIA							(
AMPLING DATES: 04/18/90-04/19/90 PA. CASE NO. 12/96 LAB: COMPUCHEM			30	tations i	01 31		cont'											
OLATILES anote ID No. raffic Report No atri nits ilution Factor ercent Moisture	NYJL-GHZ BEB19 WATER UG/L 1	NYJE-GUS DEDZO WATER Wg/L EO	NYJL-GW7 BEB24 WATER Wg/L L	NYJL-GHB BEB2S MATER Mg/L 1	PYJL - SW* BEB34 WATEP 119/1 I	NYJL-SN9 BEB3S NATER N3/L L	NTJL-SN9 DED36 MATER Ug/L I	NIJL-SMIO BER37 MATER Ug/L I	N+JL-SEE-7 BEB47 SEDIMENT Vg/Lg I S6	NTJL-5808 BEB40 SEDIMENT Ug/Fg J 46	##!!-SED? ####################################	#73L-55 0E054 SOIL ug/kg 1	NYJE-56 BEB55 SOIE vo/kg 1 23	WYJL-\$7 DE056 SOFL Ug/kg J 13	WYJL-S8 BE057 S01L Wg/kg I	MYJL-SIO DE057 SOIL Ug/19 1 35	HYJL-RIHS DEB6S WATER Wg/L I	N.Jt-RING BEDGG MATER Ug/t .l
hlorosethane ionosethane inyl Chloride hlorcethane ithilene (filcride :etone irton Disulfide 1-Dichloroethane ans-1,2-Dichloroethane libroforo 2-Dichloroethane sulanone 1,1-Trichloroethane inplicatate onodichionosethane 2-Dichlorosethane s-1,3-Dichlorosene s-1,3-Dichlorosene ichlorosethane 1,2-Trichlorosethane		110		J \$	J	J			J									j
ans-1,3-Dichleropropene poofera tethy1-2-Pentanone teranone trachioroethene luene 1,2,2-Tetrachioroethane torobenzene nylbenzene trachioroethene tene tene		3100			17	J												103537
IES: Ink space - compound analyzed for but not detected compound found in tab blank as well a sample, indicates possible/probable blank contamination estimated value estimated value estimated value, compound present below CRGL but above 10t analysis did not case EPA 04/OC Fresumitive evidence of the presence of the osterist analysis not required action lin to alayated if failution	i\$																•	be, 6

ection tin to elecated if balution

No.

ANDLING DATES: 04/18/30-04/19/90
PA CASE NO.: 13706 LAB: COMPUCHEN

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

at 10 Mo. ic Pecerl Mc. ic Pecerl Mc. bib67 bib68 bib70 Water Water Water water Water Water ug/L wg/L wg/L ug/L u	۵,
ic Perent Mc. Water Wat	
WATER WATER USTER UST USTER I	
ion factor It is is a constraint of the property of the prope	
onethane aethane (hloride oethane lene (hloride oethane lene (hloride oethane lene (hloride ichloroethane lene (hl	
osethane ethane (hloride ocethans lene Chlorist ne n n Disulfide ichloroethane ??hloroethane ichloroethane amone	
osethane oethane (hloride oethane lene Chloride ne n Disuffide (chloroethane 1.2-Dichloroethane 1.2-Dichloroethane lichloroethane nome -frichloroethane ne fichloroethane ichloroethane ichloroethane ichloroethane loroethane loroetha	
methane (hloride cethane lene (hloride cethane lene (hloride ne n	
ethane (highide cothane lene Chichide ichloroethene ?thioroethene ?thioroethene !l.2-Dichloroethene itolali plora ichloroethane anone - Irischloroethane anone ilischloroethane anone - Irischloroethane ichloroethane ichloroethane ichloroethane ichloroethane ichloroethane ilischloroethane lerechloroethane - Irischloroethane lerechloroethane - Irischloroethane - Irischloroet	
(higride options lene Chicuse ne n Disulfice (chicusethene 1.2-Dichloroethene (total) plous ichloroethane anone -frichloroethane n letrachloride neethane (chicusethane n letrachloride neethane (chicusethane chichloroethane n letrachloride neethane (chicusethane lichloroethane ne -frichloroethane ne -frichloroethane none chicusethane ne -gridelite none none none none none none none no	
oethane lene (hicust:	
lene (hlouses ne ne n Disulfide ichloroethene 11.2-Dichloroethene (total) oloro ichloroethane anone -frachloride ne ne netrachloride Acetate dichloroethane ichloroeropene loroethane -frachloroethane -frachloro	
ne n Disulfice ichloroethane -1,2-Dichloroethane lichloroethane insulfice ichloroethane anone -Irschloroethane n letrachloride Acetate dichloroeroene 3-Dichloroeroene loroethane -Irichloroethane -Irichloroethan	•
n Disulfide ichloroethane -1.2-Dichloroethane 1.2-Dichloroethane ichloroethane anone -1-rachloride n letrachloride Acetate dichleroethane ichlororoethane isoloethane loroethane loroethane -1.3-Dichloropropene loroethane en -1.3-Dichloropropene loroethane loroethane berene shoroethane -1.2-Dichloropropene loroethane berene shoroethane berene shoroethane loroethane loroetha	
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Thioreethane -1.2-Dichloroethane Italian Ichloroethane -1.2-Dichloroethane Italian Ita	
Thioreethane -1.2-Dichloroethane Italian Ichloroethane -1.2-Dichloroethane Italian Ita	
-1.2-Dichloroethene (total) plora ichloroethane anone -1richloriethane ichloroethane ichloroeropene ichloroeropene ichloroeropene ichloroeropene ichloroeropene ichloroerthane -1.3-Dichloropropene ichloroethene ince -1.3-Dichloroethane ichloroethene ince -1.3-Dichloroethane ichloroethene ince -1.3-Dichloroethene ince -1.3-Dichloroethane ichloroethene ince -1.3-Dichloroethane ichloroethane ichloroethene ince -1.3-Dichloroethane ichloroethene ince -1.3-Dichloroethene ince -1.3-Dichloroethene ince -1.3-Dichloropropene -1.3-Dichloroethene ichloroethene ic	
state of the compound analyzed for but	
ichloroethane anone -trichloroethane n letrachloride Acetate dichloroethane chloroethane -trichloropropene loroethane -trichloropropene loro -trichloropropene l	
anone -Irichlorcethane n letrachloride Acetale dichloroorooroone 3-Dichlorooroone ioroethane -Irichloroorthane -Irichloropropene lete hyl-2-Pentanone hyl-2-Pentanone chloroethane	
Itrachlorethane n letrachloride Acetale dichlereethane ichleroorethane lorethereropene lorethere lorethane litichleropropene lorethane lorethere lorethane lorethere lorethane lorethere lorethane l	
n letrachloride Acetale Acetale Acithereoethane Achteropene Boroethene Boroethane Brichloropropene Brichloro	
Acetate dichleremethane ichleremenane oroetheme icroetheme icroethane itrichleremethane itrichleremethane itrichleremethane itrichleremethane itrichleremethane itrichleremethane itrichleremenane itrichleremenan	
dichlerenethane ichlerenethane ioroethene ioroethene ioroethane ilrichlersethane ilrichlersprapene lere he ilrichlersprapene lere hyl-2-Pentanone inone chloroethene has ioroethene has ioroethene has ioroethene has ioroethane hobenzene heenzene he	
ichtoropropene 3-Dichtoropropene loropropentaine -1.3-Dichtoropropene loro -1.3-Dichtoropropene loro -1.3-Dichtoropropene loro -1.3-Pentanone	
3-Dichloropropene loroethene loroethene lorichloropropene lorichlo	
locationerthane -lrichloropropene -lrichloroprop	
space compound analyzed for but	
Trichloropropene 1.3-Dichloropropene 10:m hyl-2-Penlanone shoroethene na 2-Tetrachloroethane benzene senzene ses (Total) space compound analyzed for but	
ne -1.3-Dichloropropene lorm hyl-2-Penlanone shore thioroethene ne -2-Tetrachloroethane obenzene seenee se (Total) space compound analyzed for bul	
-1.3-Dichloropropene Crim hyl-2-Pentanone Horoethene hyl-2-Pentanone hyl-	
torm hyl-2-Pentanone chloroethene ng 2-letrachleroethane phenzene penzene penzene per compound analyzed for but	
space compound analyzed for but	
space compound analyzed for bul	·
chioroethene ng ,2-fetrachicroethane pbenzene ng servene ng servene ng compound analyzed for but	
space compound analyzed for bul	
2-letrachloroethane phenzene penzene pe penzene penze	
space compound analyzed for but	
penzene penzen	
space - compound analyzed for but	
space - compound analyzed for but	
space compound analyzed for but	
space - compound analyzed for but	
space - compound analyzed for but	
A detected	
mecunifound in lab blank as well as	

anh space - compound analyzed for but not detected - compound found in tab blank as well a sample, indicates possible/probable blank contamination estimated value estimated value, compound present below CPOL but above IDL analysis dil not pass EPA DA/RC Presunctive evidence of the tresence of the material - analysis not required ection finite elevated if Cilution

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SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

.... 02-EC. er

AMPLING DATES. 04/18/90-04/19/90
PA CASE NO.: 13/906 LAB: COMPUCHEN

romophenyl-phenyl ether

NA CHOC MATT 13.00 FMB: COUNDCHEN						(C C	nt a)								
ENI-VOLATILES	:														
ample ID No.	INT JE - GWI	NYJL-GW4	MYJL-GWS	MA) I - CMa	N. JL - GW101 DUP!	WYJL-SWI	NIJL-SW2(MS/MSD)	NYJL-SW3	NYJL-SW4	NYJL-SWA	NYJE-5WESCOUPT	MYJI -SEB2	EV31 - CCB1	1012 - 11 VII	M11-5704
raffic Report Mc.	. 00010	06021	06022	DEBZA	86827	BEB2B	8[829	BE#30	06031	8(933	BEB40	BE042	8[843	8(044	86946
atris	WATER	WATEP	WATER	WATER	WATEP	WATER	RJTAN	WATER	WATER	MATER	WATER	SEDIMENT	SEDIMENT	SEDIMENT	SEDEMENT
nits	vg/L	vg/L	ug/L	ug, t	ua/t	ug/l	wg/L	ug/L	wg/L	wg/L	ug/L	ug/kg	vg/kg	vg/kg	ug/kg
ilution Factor/GFC Cleanup (Y)	ii	i	•	i	1	i	1	i	i	i	1	1	,,, 	.99	1
ercent Meitlure		••	••	••		••	••	•••	••	••	••	24	45	40	40
***************************************		. 			· • • • • · · · • · · · · · · · · · · ·		• • • • • • • • • • • • • • • • • • • •					••••••	······	••••••	• • • • • • • • • • • • • • • • • • • •
renol	į		LAB DID	R	A										
is(2-Chloroethy) lether	:		NOT RUN		P										
Chlorophenol			ANAL YSIS	•	1										
.3-Cichlorobenzene	1		ON INIS	R	R										
.4-Dichlorobenzene	:		SAMPLE	R											
enzyt alcohot	;			•	R										
2-Dichlorobenzene	:			R	R										
Methylphenol	:			R	P										
s(2-Chioroisoprory) lether	1				R										
Methylphenol	:				R										
Mitroso-di-n-dicrocylamine	:	••		R	9							•			
rachloroethane	•			Ř											
trobenzene				8	R										
ocherene '				R	A										
Hitrophenel	•			R	R										
4-Dinethy lokenal				R											
nzoic acid					į.							1	,		3
s(2-Chloroethory inethane	į			i	i							•	- -		•
1-Dichlorophenol	•		•	ï	Ä										
2.4-Trichlorobenzene	i			R	Ř										
chialene	į			Ř	Ř										
Thiorpaniline	•				R										
<achlorobuladiene< td=""><td>1</td><td></td><td></td><td>A</td><td>P.</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></achlorobuladiene<>	1			A	P.										
.hlore-3-Methylphenol	į			R	R										
tethyinaphthalene	i			P	R										
achlorocyclopentadiene				R	R						•				
1.6-1richlorophenol	•			þ	R										
.S-Irichlorochenol	i			R	R							•			
hloronachthalene	•			R	R										
itroaniline				R	R										
ethylphthalate				R	8										
naphthy lene	i			R	R)
-Dinitrotoluene	İ			R	8										
itroaniline	•			R											
naphthene	1			•	R										3
-Cinitrophenel				R	P										
itrochenol	•			R	Ą	-							J		
entolaran	•			•	R										
·binitrotclasus	:			R	R										
thyiphthalate	i			P .	R										
plotopper/j.bpin/j. alpet	į														
Ct aus	:			Ą											J
itroaniline	•			R											
Dinitro Caethylphenol	į				P										
itrosodichenylanine	•				•										

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

SAMPLING DATES: 04/18/90-04/19/90 EPA CASE NO.: 13706 LAB: COMPUCHEN

SEMI-VOLATILES Sample 10 Mo. Traffic Recort Mo. Matrix Units Dilution Factor/SPC (Leanum (1)) Percent Moisture	HYJL-GHI BEBIB WATER Wa/L 1	NYJL-6N4 BED2J WATER Wg/L 1	NYJL-GHS BED22 Mater Wg/L	NYJL-GN9 BEB26 WATER WJ.L I	NYJL-GNIOFEUP) BEB27 MATER Ug/L I	NYJL-SNL BEB28 WATER Wg/L I	NYJL-SN2(NS/NSD) BED29 MATER Wg/L 1	NYJL-SWS DEB30 WATER Wg/L I	NYJL-SN4 BEB31 MATER ug/L L	NYJL-SM6 BER33 MATEP Wg/L t	NTJL-SM131DUP) BED40 MATEP Ug/L I	NYJL-SED2 BEB42 SEDIMENT Ug/kg 1 24	NTJL-SED3 BED43 SEDIMENT Ug/kg 1 45	NYJL-SED4 BEB44 SEDIMENT Ug/kg . 39 40	NYJL-SED6 BED46 SEDIMENT Ug/kg L 40
Pentachlorophenol	1	•••••	•••••	R	R	• • • • • • • • •	••••••	•••••	••••••	• • • • • • • • •		••••••••	*********	••••••	**********
Phenanthrene	:			R	#										J
inthracene	:			R	R										3
Di-m-tutylohth-late	!			f											J
lvoranthene				R	R										1400
) y i ene	į			ı.										1	1200
iutylbenzyšphthalate	į				×										7
1.1'-Pichlorobenzifine -enzolalanthiacene	1											111			818
hrysene	-			ì								3N 3N :			810 870
is(2-Elhytheryt)chthalate	;	••		ï	ì							## :		580	3000
i-n-octylphthalate	i			Ř	ä										1
enzoibifluoranthene					R							JH	JW	TI,	2000 Ex
enzol h If I war antheire	:				A							JH	JH	JN	2000 EN
entol a juli sue	!			R	R										760
ndeno[1,2,3-cd]pyrene	:			R	•										j
ibenzfa,h)anthrace ne	:			R	R										J
anzolg.h.i Ivervlene	:			R	Ř										J

0165:

lamb scace · compound analyzed for but not detected

- compound found in lab tlank as well as sample, indicates possible/probable blank contamination
- estimated value
- estimated value, compound present below (ROL but above IDL
- analysis did not pass (PA QA/QC
- Presumptive evidence of the presence of the material
- analysis not required

tection limits elevated if Dilution ctor of and/or percent moisture >0%

10354

Der. C.

SUbmary Ur sile inspection ANALTICAL DATA (cont'd)

ELM CH2C HAT. 1240. FMB: COULDENE						(cont'd	1)				
SEMI-VOLATILES	;										
Sample 10 Mo.	!R-JCYN!	NYJL-SZIMS/MSD)		MY36-24	MA76-23	HYJL-511109P)	MYJL-RIMI		NYJL-AINS		
Traffic Pencil Mo	 B{B50	PEBS1	EEB52	86852	86836	26860	BEBEI	06065	86863	86864	86847
Matri.	; 501t	SOIL	SOIL	501L	SOIL	SOIL	MUIEs	MATER	WATER	WATER	WATEP
Units	: wa/ka	nė/r>	ua/la	uā/l à	va/) g	ug/kg	ug/l	wg/L	ug/L	vg/t	vg/L
Dilution Factor/SPC Cleanup (Y)	; ι	. 39	l	1	1	1	1	1	1	ı	M/A
Percent Moisture	; 27	52	16	12	12	24	••	••	••	••	M/A
Phenol	······································	• • • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • •	• • • • • • • • • •				••••••	•••••		#R
bist2-Chlorcethyl lether	į										HP
2-Chlorochenol											MŘ
1.2-Dichlorobenzene	;										MR
1.4-Dishlorobentene	•										MR
Benzyl alcohol	:										MR
1.2-bichlorotenzene	:										M2
2-Bethylchenol	:										KA
oist2-Chloroisonropy1 lether	•			•							KR
											NA NA
1-Methylphenol											
-Mitroso-di-n-dipropy lamine	į										NR NA
lexachleroethane	•	••									MA
litropentens	i										MA
Isophorone	•										押
!-Nitrophene! .	į										199
?,4-Cinethylchenol	;										MR
rentoic acid	:	1	3		J	3					WR
is(2-Chloroethoxy)nethane	:										KR
.4-Dichlorephenol	:										眦
.2.4-Trichlerebenzene	1										MR
aphthalene	•										NA.
-Chloroaniline											MR
exachlorotutadiene	i										NR '
-Chlore-3-Methylphenol	i										MR
-Methylnaphthalene	į										KR
exachlorocyclopentadiene	į										MR
.4.6-Trichlorophenol											MA
.4.3-Trichlorophenol											NR.
·Chloronaphthalene	•										MR
-Nitroaniline											nr
inethylphthalate	:				_						WR
cenaphthylene	;				•						MR
,6-binitrotoluene	:										WR
-Nitroaniline	;										WR
	:										NA.
senaphthene	:										FR
.4-Dinitrophenel	•										NA
Mitropheno!	į										NP
ipeutojai au	:					•					AP
.4-binitrotolvene	:										RA MA
iethylphthalate	:										MR
Chlorophenyl-chenyl ether	:										MF
luotene	:										HP
Mitroaniling	:										HR
6-Cinitro-2-methylphenol	;										NP
nitrosodichenylamine	•										MR
Branghenyl-chenyl ether											NP
reach) or observers	•										PP

EPA CASE NO .: 13709 LAB: COMPUCHEM

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SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

SEMI-VOLATILES Sample ID Mo. Traffic Recort Mo. Hatrix Units Dilution Factor/GPC Cleanup (Y) Percent Moisture	NYJL-51 BEB50 SOIL ug/kg I 27	NYJL-S2(MS/MSD) BEB51 S01L wg/kg . 99 52	NYJL-\$3 DEB\$2 \$01L ug/kg 1 1B	NYJL-\$4 6(053 501L ug/kg 1	NYJL-59 BEB58 SOIL Ug/kg I 12	MYJE-SLEEDUP BEB60 SOIL wg/kg I 24	NYJL-RIN1 DED61 WATER Ug/L I	NYJL-RENZ BE862 MATER Wg/L I	NYJL-RINS BEB63 MATER Wg/L I	NYJL-REN4 BED64 MATER Ug/L I	NYJL-TROK] BEB69 MATER Ug/L H/A M/A
Pentachlorophenol											NA
Phenanthrene	:				3	3					WR
Anthracene		_			J						MR
Di-n-butylphthalate	1	,	630								MR
Fluoranthene					610	,					yr
Pyrene	•	•			530	,					KA Ha
Butylbenzylphthalate 3,3'-Dichlorobenzidine											NP. NP
genzola Janthracene	:				410	,					HR HR
Chrysene Chrysene					370	,					MR
bis(7-Ethylhenyl)phthalate	; ,	•	,		374	1					WR
Di-n-ectylphthalate	•	. . •	•			•					KR
Denzo(b)/luoranthene	: JN	Jn			780 E	M 460 E	n				NR.
Denzo(k)fluoranthene	311	JK			780 E						##
Benzola)pyrane ·		***			440	,					WA.
Indenoi 1,2,3-cd byrene		•			1	j					MR
Dibens (a. h)anthracene	į				1						MR
Benzola, h. i Iperylene					380	J					MR

NOTES:

Blank space - compound analyzed for but not detected

- B compound found in lab blank as well as sample, indicates possible/probable blank contamination
- [· estimated value
- J estimated value, compound present below CROL but above 1DE
- ₹ analysis did not pass EPA QA/QC
- Presumptive evidence of the presence of the material
- IR analysis not required

Detection limits elevated if Dilution actor >1 and/or percent moisture >0%

SAMPLING DATES: G4/19/90-04/19/90
EPA CASE NO.: 12704 LAB: COMPUCHEM

lexachloroben:ene

TABLE 1 SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

			301	, , , , , , , , , , , , , , , , , , ,	<i>.</i> 011	(c	ont'd))									•	
SEMI-VOLATILES						-	-		W. 15. 875.	W-41 4864	W. M. *F&A							
Sample ID No.										PYJL-SCD9			NYJL-\$6	MYJL-57	NYJŁ-50	NYJŁ-510		
Traffic Report No.	; 00019	06950	DEB24	86652	06834	DE035	BE036	BEB37	BEB47	8848	8(849	BED54	BEBSS	BED56	BEB57	88859	86865	DE044
Hatris	HATER	WATER	WATER	WATER	WATER	MATER	WATER	WATEP	SEDIMENT	SEDIMENT	SEDIMENT	SOIL	2016	SOIL	SOIL	SOIL	WATER	WATER
Units	wg/L	ug/L	wg/L	ug/l	wg/L	ug/L	wg/L	wg/L	wg/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	ug/kg	wg/L	ug/L
Dilution Factor/GPC Cleanup (Y) Percent Noislure		 					! 		2 56	146	83	1 18	1 23	13	t 18	1 35	 	
Pheno)		R	,		••••••	R	A	•••••	j			•••••	••••••	• • • • • • • • • • • • • • • • • • • •	,	• • • • • • • • •		
bis(2-Chloraeth, 1 lether	;		•															
2-Chlorophenol	:	R				R	R											
1,3-Dichlorobenzene	:																	
t.4-Dichlarabenzene	:																	
Benzyl alcohol	:						•											
1.2-Dichlorobensene	:																	
2-Hethylphenol	:	R				R												
bis(2-Chloroisecropyl)ether	:																	
4-Hethylphenol	:	R				*	R											
N-Mitroso-di-n-diprocylamine	•													:				
Hexachloroethane		••																
Hilrobenzene	i																	
Isochorone	•																	
2-Mitrophenol	i	R				R												
2.4-Dimethylchenol	i	R				R	R											
Dentoic acid	i	Ř					P	J	J							3 1		
bist2-Chloroethoxy Inethane	i	••						-								-		
2.4-Dichlerephenol	i	ı.				£	R											
1,2.4-Trichlorobenzene	1											J						
Naphthalene												·		j				
4-Chloroaniline																		
Hexachlorotutadiene	į																	
4-Chloro-3-Methylphenol	;				3		R											
7-Methylmaphthalene					-													
Mexachlorocyclopentadiene																		
2.4.6-Trichlorophenol		R					R											
2.4.5-11 ichiorophenol		ï				Ř	Ř											
2-Chloronachthalene		•				-	•											103543
2-Mitrosniline	•																	U
Disethylphthalate	•												•					44
tenachihylene	;																	ω
2,6-Danitrololuene	:																	
3-Mitrosnijine	;																	
Jeuschfpeue	;								J					j				
	:					R			•					•				
2.4-Dinitrophenol	;						Ř											
I-Nitroshenol		K				•	•		J					,				
ibenzoluran									,					•				ייב
2,4-Cinitrotoluene	į																	Rev.
rieth, lohthalate	•												•				_	
i-Chlorophen.1 ohenv1 ether	•								1					j			ھ	
luorene									,					,			~	70 8
1-Milroaniling		_				•											~	
1.e-Dinitro-2-eethylphenol		R				A	•										_	·'' o
i-mitrosodichenylamine	•																84 3°18	6
1-Bromophenyl-phenyl ether	:																e)	, 0,
In-rabins at an ease	•																	

SAMPLENS THATES: 04/18/30-04/19/90 EPA FASE NO.: 12706 LAB: COMPUCHEN

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

SEMI-VOLATILES Sample ID Mo. Traffic Recort Mc Matric Units Dilution Factor/SPC Cleanum (Y)	HYJL-GN2 BEB19 MATER Ug/L	NYJL-GN3 BEB20 MATER Mg/L 1	NYJL-GN7 BEB24 WATER Wg/L	NYJL-GN8 BEB25 MATEP Ug/L L	HYJL -SW7 DEB34 WATER WG/L L	NYJL-SH8 BEB35 MATER Ug/L L	NYJL-SW9 BEB36 WATER Ug/L 1	NYJL-SWIO BEB37 WATER Ug/L 1	NTJL-SED7 BEB47 SEDIMENT Ug/kg 2	NYJL-SED8 BEB4B SEDIMENT Ug/kg I	N/JL-SED9 BEB49 SEDIMENT ug/kg I	NYJL-SS BEB54 SOIL ug/kg 1	HYJL-56 BEBSS SOIL ug/kg 1	MYJL-37 BEB56 SOIL wg/kg 1	MYJL-SB BEBS7 SOIL Ug/kg 1	MYJL-S10 BEB59 SOIL ug/kg 1	NYJL-RINS BEB6S WATER Wg/L 1	NYJL-RING BEBGG WATER Wg/L 1
Forcent Hossture	•	••	••	••	••	••	••	••.	56	46	83	18	23	13	18	35	••	••
Fentachlorophenol		•••••	••••••	• • • • • • • • •	• • • • • • • • • •		•	*******	• • • • • • • • • • • • • • • • • • • •	········	•••••••	••••••	•••••	•••••	••••••	••••••	*	••••••
Phenanthrene	:	•				-			2700			j	J	890		1		
Anthracene	į								J				-	J.		•		
Di-n-buty);hthalate	İ								J							J		
Fluoranthene	!								6830			J	1	970		J		
Parene	1								3900			J)	250		3		
Butyltenzylphthalate									J					J		J		•
3.3'-Cichlarobenzidine																		
Benedia Fanthracene									2400				3	560		j		
Chrysene									3000			J	,	540	_	j		
bisi2-Ethylhenyljohthalate	; 1	••							7290			,	J	:)	,	,		
Di-n-octylohthalate BenzolbJf Iworanthene	i i			٠.					, 3600			111	JW	130 (?m	14		
Benzolk If I wor anthene)								2500			JN	7# 1#	730 (JM 16		
Bento(a)bytene •									2500			3)]	420	r	3"		
Indensi I. 2.3-cd byrene									1			•	j	1		j		
(ibenzia, h)anthracene									j				•	j		•		
Benzolg.h.i Iservlene									j				J	j		J		

HOTES:

Blank space - compound analyzed for but not detected

- 8 compound found in lab blank as well as sample, indicates possible/probable blank contamination
- E estimated value
- J estimated value, compound present below CPOL but above IDL
- 4 analysis did not pass EFA QA/OC
- + : Presumptive evidence of the presence of the material
- ranalysis not required
- Detection limits elevated if Dilution
- factor of and/or percent moisture >O1

rachloroteriene

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

EHI-VOLATILES	;		
anole ID Ho.			NYJL-TROKZ
raffic Recort Ho.	1 01067	868	BER70
latrix	WATER	WATER	WATER
inits	ug/L	wg/L	ug/L
ilution factor/GPC Cleanup (Y)	1		#/A
ercant Moisture			n/a
henol			MR
isl2-Chlorgethyllether	:		MA
·Chlorochenol	:		WP.
.3-Dichlorotenzene	;		MR
.4-Dichtorotenzene			MR
entyl alcohol	:		MB
.2-Dichlorotenzene			MR
-Methylphencl	•		ME
ist2-Chloroisopropy1 lether	i		MR
-Nethylphenol			NP.
·Nitroso-di-n-dicropylamine	i	••	NF
ewachloroethans	i		NA
itrobenzene	i		11P. 14P
sopher one	i		117 118
-Mitrochenol .4-Dinethylchenol	•		
entoic acid	· •		DŽ
is(2-Chloroethoxy)methane	:		
.4-Dichlorophenol	i		***
.2.4-Irichlaratene	ļ		MR
iohthalene			NR
Chloroaniline	•		NR
exachlorobutadiene	i		118
Chlora-3-Methy Johenai			MR
Mathylmachthalene			NP
:machlorocyclopentadiene			MR
4.4-Trichlorophenol	i		MP.
4.5-Trichlorophenol			MA
Chloronaphthalene	:		MR
Mi,troaniline	;		MR
methylphthalate	:		神界
enachthylene	•		KP
6-Dinitroteluene			MR
Mitroaniline			MR
enaphthene			MR
4-Dinitrophenal			NE
Mitrochenol			NR
benzoluran			NB WA
4-Pinitrotolvene			MR
ethyiphthaiate	:		MR
(hlorophen.1-phenyl ether			NS.
uorene			NA ***
Mitropoiline			HP
6-finitro-2-methylphenol	•		MA
nifrosodiphenylamine	į		#R
Broophenyl-chenyl ether	i		AN De
1 15 3 1 0 1 0 1 0 1 0 1 0 0 0 0 0 0 0 0 0 0	•		26

Rev

P.F. 6

SAMPLING DATES: 04/18/70-04/19/70 EPA CASE NO.: 13706 LAB: COMPUCHEN

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

SEMI-VOLATILES Sample ID Mo. Traffic Recort Mo. Matrix Units Dilution Factor/GFC Cleanue fri Percent Moisture	BEB67 MATER Ug/L 1	NYJL-RENZ DEB4B WATER Wg/L I	NYJE-TRBF2 BEB70 WATER Wg/L H/A H/A
Pentachlorochenol		•••••	NR
Phenanthrene	:		MP
Anthracene			MR
(i-n-butylehthalate	:		MR
Luoranthene	:		NR
Pyrene	;		MR
Butylbenzylohthalate			MR
3.3'-Dichlorobentidine			MR
Benzola Banthir acene			MR
Thi ysene	į		MR
ois(2-Ethylheyyl)ohthalate	į		NR NA
ri-n-octylphthalate	•	••	MR.
renzolb)/I wor anthene	•		MA MA
leuto(s)chiene .	;		118
Indenoil, 2, 3-cd loyrene	;		
ribensta.hlanthracene			12
enzolg.h.i)cerviene			MR

01ES:

lank space - concound analyzed for but not detected

- compound found in lab blank as well as sample, indicates possible/probable blank contamination
- · estimated value
- estimated value, compound present below CROL but above IDL
- analysis did not pass EPA QA/QC
- Presumptive evidence of the presence of the material

A - analysis not required

election limits elevated if Dilution actor of and/or percent moisture 'Ot

103546

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

ole ID Mo. ffic Recort Mo. ri> ts ution factor/GPC Cleanup (Y) cent Moisture	RYJL-GNI DEDIB WATER Wg/L	NYJE-5N4 OEB21 WATER Ug/L I	RTJL-GNS BEB22 WATER Ug/L I	BEB26 WATER Wg/L L	BED27 Water Ug/L I	DEB2B WATER Ug/L I	NYJL-SN21NS/NSD BEB29 WATER Ug/L I	BEB30 WATER wg/L J	BEB31 WATER Ug/L 1	BEB33 WATER Ug/L I	BED40 WATER Ug/L 1	BE042 SEDIMENT Ug/kg I 24	BEB43 SEDIMENT Ug/kg 4 45	BEB44 SEDIMENT ug/kg 1 40	BEB46 SEDINEN Ug/kg I 40
ha-BMC a-BMC ta-BMC ta-BMC (Lindane) tachlor rin tachlor eroxide osulfan 1 drin			R R R R R	•••••				•••••	•••••			,			R R R R
-COE in sulfan II -OCO sulfan sylfate -COI pwychlor												;			
n ketone -Chlordane -Chlordane hene or-1016															
or - 1221 or - 1232 or - 1242 or - 1248 ór - 1254 or - 1260						2		7.6 1.4			2.2	7600 1600	50090	610	

As space - coopound analyzed for but not detected coopound found in lab blank as well as sample, indicates possible/probable blank contamination estimated value estimated value compound present below CRQL but above IDL analysis did not pass EFA QA/QC Presunctive evidence of the presence of the material analysis not required ction limits elevated if Dilution or 11 and/or percent moisture 'Ot

Period

1000: 02-9003-01 SUMMARY OF SITE INSPECTION ANALYTICAL DATA SAMPLING DATES: 04/18/90-04/19/90 (cont'd) EPA CASE NO : 13909 LAB: COMPUCHEN

aloha-BMC beta-BMC delta-IMC delta-I	PESTICIDES Sample ID Mo. Traffic Report Mo. Hatrix Units Dilution Factor/GPC Cleanup (T) Percent Moisture	NYJL-S1 BEB50 SOIL ug/kg I 27	NYJL-52(MS/MSD) DED51 SOIL .ug/kg .1 .52	NYJL-S3 BEB52 SOIL ug/kg I	NYJL-S4 0E053 SOIL ug/Lg 1	NYJL-59 BEB58 SOIL Ug/kg I	MYJL-SII(DUP) BEB60 SOIL ug/kg I 24	NYJL-RINS DEB61 WATER Ug/L I	NYJL-RINZ BE062 WATER Wg/L I	NYJL-RINS DED63 WATER Wg/L 1	NYJL-RINA BEB64 WATER Ug/L 1	NYJL-TABKI BEB69 MATER Wg/L M/A M/A
RE RE RE RE RE RE RE RE			••••••		• • • • • • • • • •	• • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •	•••••	•••••	•••••	•••••••	
R		i		K								
Rectachior		•		R B								
Aldrin R R R R R R R R R R R R R R R R R R R		•		ï								
Endosulian I R Dieldrin 4.4-DDE 34 HR Endrin Endosulian II HR 4.4-DDE HR Endosulian II HR 4.4-DDE HR Endosulian II HR 4.4-DDE HR Endosulian Sulfate 4.4-DDE HR Endosulian Sulfate 4.4-DDE HR Endosulian HR 4.4-DDE HR Endosulian H		i		 R								
Dieldrin	Heptachlor evoxide	•		*								MR
4.4'-DDE		+		R								WR
Endrin		1.										•
Endosulfan il 4,4'-000 MR Endosulfan sulfate MR 4,4'-001 60 57 MR Hethosychior MR Endrin ketone MR alpha-Chlordane MR ganna-Chlordane MR froctor-101c MR Aroctor-1221 MR Aroctor-1221 MR Aroctor-1232 MR Aroctor-1242 MR Aroctor-1248 #R		34										****
4.4'-000 Endosulfan sulfale 4.4'-001 60 57 IR Hethosychlor Endrin ketone alpha-Chlordane ganna-Chlordane floxaphene Aroclor-101c Aroclor-1221 Aroclor-1221 Aroclor-1242 Aroclor-1248 AR HR			••									••••
Endosulfan sulfate 4.4'-bbī 60 57 MR Hethosychtor Endrin ketone alpha-Chlordane ganna-Chlordane froclor-10j£ Aroclor-1221 Aroclor-1221 Aroclor-1232 Aroclor-1242 Aroclor-1248 ###################################												•
4.4°-DDT 60 57 MR Methosychlor Endrin ketone Alpha-Chlordane Aganna-Chlordane Aroclor-1012 Aroclor-1221 Aroclor-1232 Aroclor-1242 Aroclor-1248 Aroclor-1248	,	į										
Methosychior MR Endrin ketone MR alpha-Chlordane MR gamaa-Chlordane MR Toxaphene MR Aroclor-101z MR 4roclor-1221 MR Aroclor-1232 MR Aroclor-1242 MR Aroclor-1248 MR		4.6					63					
Endrin ketone alpha-Chlordane gamaa-Chlordane foxaphene Aroclor-10jt Aroclor-1221 Aroclor-1232 Aroclor-1242 Aroclor-1248 Aroclor-1248		;					37					****
alpha-Chlordane MR gamaa-Chlordane MR Toxaphene MR 4roclor-10Jz MR 4roclor-1221 MR 4roclor-1232 MR 4roclor-1242 MR 4roclor-1248 4700 MR												
Toxaphene		i										MR
Aroclor-1016 Aroclor-1221 Aroclor-1232 Aroclor-1242 Aroclor-1248 Aroclor-1248 Aroclor-1248 Aroclor-1248 Aroclor-1248 Aroclor-1248	gamma-Chlordane		′									MA
4roclor-1271 NR 4roclor-1232 NR 4roclor-1242 NR 4roclor-1248 NR	Toxaphene	:										MR
Aroclor-1232 MR Aroclor-1242 MR Aroclor-1248 A700 MR	Aroclor-101e	:										MR
Aroclor-1242 Aroclor-1248 4700 MR		;										MR
\roc1or-1248 4700 NR												
,		•										•••
2700		i	•									
rroclor-1266 NR						7700						

101ES:

lank space - compound analyzed for but not detected

- I compound found in lab blank as well as sample, indicates possible/probable blank contamination
- · estimated value
- 1 estimated value, compound present below CPOL but above IDL
- ! analysis did not pass EPA QA/QC
- · Presumptive evidence of the presence of the material
- R analysis not required
- etection limits elevated if Dilution
- actor of and/or percent moisture :0%

TICIDES NYJL-SUB NYJL-SU7 NYJL-SUB NYJL-SU9 NYJL-SU10 NYJL-SED7 NYJL-SED8 NYJL-SED9 NYJL-SS ple ID No. HYJL-56 MYJL-S7 MYJL-SB MYJL-SIG MYJL-RIMS MYJL-RIM6 !NYJL-GW2 NYJL-GW3 NYJL-GW7 ffic Report No. : BEB19 BE820 BEB24 **BEB25 BEB34** BE 835 36936 DEB37 **BEB47** 86848 BE849 BEB54 86855 16856 38857 BEBST WATER WATER WATER WATER SEDIMENT SEDIMENT SEDIMENT SOIL SOIL WATER WATER WATER WATER SOIL SOIL SOIL wg/L ug/L wg/L vg/L ug/L ug/L ug/L wg/L ug/kg ug/kg ug/kg ug/kg ug/kg ug/kg ug/ke wq/kq ution Factor/GPC Cleanup (Y) 1 t 1 1 1 1 1 1 1 ı 1 1 1 46 83 cent Moisture 56 18 23 13 18 35 ha-BMC ta-BHC ma-BHC (Lindame) tachlor tachlor eposide 67 E osullan 1 170 sulfan II 0.17 79 150 osulfan sulfate 71 wxychler. in betone ia-(h)ordane ia-Chlordane ohene :lor-1016 Jor-1221 :lor-1232 1or-1242

540

640

430

TABLE 1

SUMMARY OF SITE INSPECTION ANALYTICAL DATA

(cont'd)

S: k space - compound analyzed for but not detected compound found in lab blank as well as sample, indicates possible/probable blank contamination estimated value estimated value, compound present below CROL but above IDL analysis did not pass EPA QA/QC Presupritive evidence of the presence of the material analysis not required ction limits elevated if Dilution or >1 and/or percent moisture >0%

F: 02-7003-01

ris

ls

3-EMC

rin

Idrin .-00E

in

-000

100

1or-1248 lor-1254

lor - 1260

PLING DATES: 04/18/90-04/19/90

CASE NO : 13706 LAB: COMPUCHEN

02-9003-01 Rev. No. (

01065

WATER

we/L

1

16846

WATER

wq/L

1

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

SAMPLING DATES: 04/18/90-04/19/90 EPA CASE NO.: 13906 LAB: COMPUCHEN

PESTICIDES Sample 10 Mo. Traffic Report Mc. Hatrix Units Dilution Factor/GPC Cleanup (Y) Percent Moisture	NYJL-RIN7 BEB67 MATER Wg/L 1	NYJL-RIMB DED68 MATER Wg/L 1	NYJL-TRBFZ BEB70 WATER Wg/L M/A M/A
alpha-BHC beta-BHC delta-BHC delta-BHC delta-BHC qanna-BHC (Lindane) Heptachlor Aldrin Heptachlor eroxide Endosulfan 1 Dieldrin 4.4'-DUE Endrin Endosulfan 11 4.4'-DUO Endosulfan sulfate 4.4'-DUO Hethoxychlor Endrin tetone alpha-Chlordane qanna-Chlordane Toxabhene Aroclor-1016 Aroclor-1221 Aroclor-1242 Aroclor-1248 Aroclor-1254 Aroclor-1254			神界 神泉 神泉 神泉 神泉 神泉 神泉 神泉 神泉 神泉 神泉 神泉 神泉 神泉

NOTES:

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Blank space - compound analyzed for but not detected

- B compound found in lab blank as well as sample, indicates possible/probable blank contamination
- [· estimated value
- J estimated value, compound present below CROL but above IDL
- R analysis did not pass EPA QA/QC
- M Presumptive evidence of the presence of the material
- MR analysis not required
- Detection limits elevated if Dilution
- factor 1 and/or percent moisture -0%

TE MAME: LI TUNGSTEN DB: 02-7003-01

IMPLING DATES: 04/18/90-04/19/90

'A CASE NO.: 13906 -B NAME: ENSECO/RMAL

TABLE 1 SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

ORGANICS mple ID Mo. affic Report Mo. triv its	NYJL-GHL MBCJOI WATER UG/L	NYJL-GH4 NDCJO4 NATER Ug/L	NYJL-GNS NBCJOS WATER Wg/L	NYJL-GN9 MBCJ09 WATER Ug/L	NYJL-GH10(DUP) H8CJ10 HATER UG/L	NYJL-SUL HBCJII HATER Ug/L	NYJL-SM2(MS/MSD) MBCJ12 WATER Ug/L	NYJL-SN3 MBCJI3 WATER Ug/L	NYJL-SW4 MBCJ14 WATER Ug/L	NYJL-SN6 MBCJ16 NATER Ug/L	NYJE-SWES(DUP) MBCJZS WATER WG/E	NYJL-SED2 MBCJ25 SEDIHENT mg/kg	NYJL-SED3 MBCJ26 SEDIMENT ag/kg	NYJL-SED4 MBCJ27 SEDIMENT Bg/Fg	NYJL-SEO6 MBCJ29 SEDIMENI mg/kg
uĐi nuh	32600	171000		122000	120000	j	409	931	204	j	J	2190	3560	12600	3 6009
timony	68.8		3390	184	212		73.1	212				3320 E	3290 E	245 E	
senic	1	J	R	2690	2800	J	50.2	145 E	15.2		j	1240	1770	228	20.1 €
rium	525	707	,	J	J	1	J	j	J	1	J	333	387	131	3
ryllium	1 1	11.1		11.2	12.3									2.8	3
deive												5.6	10.1	5.9	€.3 €
lcium	26100	28000	214000	541000	572000	6560	6670	40600	449000	71200	6930	2800	24000	149000	4440 E
roniun	97.9	344	3	137	132		, , ,	,,,,				24.5	78.7	62	34.5 E
balt	128	221		353	358		61.2	475	85.8			18.1	60.8	1390	53.6 E
pper	171			231 8		J	103 E		48.2 E	J	26.4	. 171 E		994 E	
3 n	228000	257000	6390	370000	384000	.174	2150	4530	547	871	, 141	46400	145000	20900	19200 E
3 d	31.1	209	Ę	144 [•	, 141	195	102 E	175444	•	2950	5140	937	254 E
gnesium	17900	46300	J	179000	187000	•	J	,,,,	15900	175000	•	, 221	, 260	24100	4240 E
nganese	3990	7620	1730	35300	37300	•	100	535	138	173	•		• • • •	1110	245 E
rcury	0.42	13	1.41	0.28	0.25		0.21	0.66			•	4.4 E	9 E	0.71 E	
:kel	135	213		339	336	•	, 50	140	, 1000 t	47000 1		•	3430	271 3510	\$2.3 E
tassiun	10100	E 14100 E	•	25400 E	25600 E	,	•	· ·	5920 E	17000 (. •	•	3430 19 E	231A	•
leniva	İ					•	15	23.6	•			103	136	32.2	33.8 E
iver tius	10600	7940	13100000	1390000	1460000	36100	14100	56300	21700	1360000	36700	144	1	32.2	5150 E
illius	1 10000	7790	13100000	137000	140000	30100	14100	30300	21700	130000	30700	•	•		3130 €
vadium	125	512		198	181							•	•	56.5 €	28.8 E
10	508		. 1	5940 E		20.7 E	93.3 E	229 E	31.4 E	3	R	119 E	197 E		
ınide	1 ,00	13.6	•	3,44 6			70.0 0	••••	J1	•	•	••••	• •		
111145	1	17.7													

ES:
.nk space - compound analyzed for but
not detected
estimated value
estimated value, compound present
below CRUL but above IDL
analysis did not pass EPA QA/QC
- analysis not required

103551

265.6

SITE NAME: LI TUNGSTER

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

SAMPLING DATES: 04/18/90-04/19/90

EPA CASE NO.: 13906 LAB HAME: EMSECO/RHAL

INDRGANICS Sample ID Mo. Traffic Report Mo. Matrix Units	MYJL-S1 MBCJ33 SOIL mg/kg	MTJL-S2(MS/MSD) MBCJ34 S01L ag/kg	NYJL-S3 MBCJ3S SOIL mg/kg	NYJL-S4 MBCJ36 SOIL mg/kg	NYJL-S9 MBCJ41 SOIL ng/kg	MYJL-\$11(DUP) MBCJ43 S01L mg/kg	NYJL-RINI MBCJ44 Water Ug/L	NYJL-RIN2 MBCJ4S WATER Ug/L	NYJL-RINS MBCJ46 MATER Vg/L	NYJL-RIN4 MBCJ47 WATER Wg/L	NYJL-TRBRT N/A N/A Ug/L
liuminum	6150	32300 €	330	519	4960	5550			•	•	MR
\ntimony	258 E	272 E	796 E	188 E	296 E	189 €					HR
rsenic	309	2600 E	3370	3700	233	413					MR
Barium	į į	492 €	J	J	177	J					MR
teryllium	,	8.9 E		•	J						MR
ladejus	; ;	49 E	14.9	16	9.1	1.5					WR
alcium	, ,	59200 E	J	J	2200	3					MA
Thronium	14.3	172 €		39.1	20.5	14					MR
obalt	; ;	198 E	3	J	73.5	J					NR
opper	46.2 E	30€0 €	2190 E	752 E	1150 E	46.5 E			J	J	MR
ron ·	28500	"172000 E	327000	246000	60100	31600	1	J	1		HŘ
ead	179	16000 E	9090	1960	8660	198					MR
lagnesiwa	; ;	2470 E	3	J	1540	j					MR
langanese ,	87.3	25100 E	2260	52 90	2120	85.4					MR
ercury	0.54 E	13 E	0.48 E	0.6 E	1.4 E	0.37 E					糖
ickel	; ;	53.9 E			165	j					ĦR
otassium	; ;	J	J	J	J	J					HR
eleniva	:		2.5 €	3.4 E	10 E						MR
ilver	1.8	65.5 E	156	84.3	55.5	5.5					MR
odive	:		8540			J					MR
hallium		R	R	R	ŧ	R					HR
anadium	20.8 E		1	3	22.7 €	20.3 E					HR
inc	43.3 E	2980 E	1330 E	1240 E	559 E	39.7 €				J	ĦŘ
yanide	:	1.5 E									MR

ITES:

lank space - compound analyzed for but

not detected

- estimated value

- estimated value, compound present below CRDL but above IDL

- analysis did not pass EPA QA/QC

1 - analysis not required

103552

R.F. 6

TTE WAME: LT TUMGSTEN DDT: 02-9003-01

AMPLING DATES: 04/18/90-04/19/90

PA CASE NO.: 13906 AB NAME: EMSSCO/RMAL

TABLE 1 SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

MORGANICS ample 1D Mo. raffic Report Mo. atri> nits	NYJL-GN2 HBCJ02 NATER UG/L	NYJL-GNS MBCJO3 MATER Ug/L	NYJL-GW7 MBCJO7 WATER Ug/L	MYJL-GWB MBCJOB WATEF Ug/L	NTJL-SN7 MBCJ17 MATER Wg/L	NYJL-SN8 MBCJ18 MATER Ug/L	NYJL-SN9 MBCJ19 WATER Ug/L	NTJL-SHIO MBCJZO WATER Ug/L	NYJL-SED7 MBCJ30 SEDIMENI Mg/Lg	NYJL-SED8 MBCJ31 SEDIMEHT mg/kg	NYJU-SED9 PBCJ32 SEDIHENT mg/kg	WiJL-SS MBCJ37 SOIL mg/kg
lyninyn	152000	115000	3970	3710	j	581	338	J	9200 E	7570	5:40 E	16300
ntimony	į ,		276	J		86.2		817			104 E	316 E
rsenic	, ,	J	\$1.7	31.8		43.8	17.2	J	17.1 E	45.7	193 E	437
arium	963	874	j	1	1	3	;	J	J	196	1340 E	860
eryllium	11	3.4							J	J	J	2
adeiun	53.6	14 E	29.1		•	7.5 €		15.2 E	R		J	16.5
alcium	39900	88100	90600	130000	32400	53500	37700	37500	19960 E	3	13200 €	36800
hronius	369	271	23.7	20.9				66.2	47.4 €	20.4	87.6 E	160
tale	; 115	. 109	703	52.9		:320	305	17500	30 E	92.1	7910 E	2270
19990	; 231 E	2080 E	391 8	60.8 E	J	167 E	55.9 E	1570	368 E	26.1 €	571 E	4180 E
ron	246000	231000	9450	34200	663	17300	24200	6580	21000 E	22000	237000 E	85900
ead	146	121	198	7.3		30.8	13.2 €	153	345 E	31.5 E	354 €	3390
agnesiwa .	57200	42500	13000	19300	21000	14700	12700	18400 €	7990 E	1990	1	48700
Inganese	3900	8190	1480	829	93.5	7780	7200	27500	515 E	205	65100 E	18400
:rcury	0.25		0.87	0.29				0.84	0.53 E			1.7 €
ickel	243	225	525	50		1700	367	76200	40.3 E	44.4	33:0 E	9130
itassium	; 34100 E	11490 E	8720 E	9380 E	7290 E	8370 E	J	7633	J	J	J	3
:leniu o	:		J									1.9 E
lver	:		J					34.6	37.6 E		23.6 E	75.6
ndium	; 23300	45500	117000	35900	202000	67800	21300	464000	13500 E			4460
nallium									R	8	Ř	R
nadium	421	320	J	J		J			37.7 €	32.9 E		43.3 E
nc	\$ 632 E	1820 8	3840 E	160 E	1	1530 E	81.4 E	,	300 [96.9 E	622 E	1570 E
anide	:	11.4										

IES:

ank space - compound analyzed for but

- not detected
- estimated value
- estimated value, compound present below CRDL but above 10t
- analysis did not pass EPA QA/QC
- · analysis not required

T03553

125,6

ITE NAME: LI TUNGSTEN DDD: 02-9003-01 TABLE 1
SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

HAMPLING DATES: 04/18/70-04/19/90
TPA CASE NO.: 13906

TPA CASE NO.: 13906
AB NAME: EMSECO/RMAL

INDRGANICS Tample ID Mc. raffic Report Mo. fatrix Init:	HYJL-S6 HBCJ39 SOIL Bg/kg	NYJL-57 MBCJ39 SOIL ag/kg	MYJL-S8 MBC340 SOIL ag/kg	NTJL-S10 MBCJ42 SOIL mg/kg	NYJL-RINS HBCJ48 HATER UG/L	HYJL-RING MBCJ49 WAIER Wg/L	NYJE-RIN7 HBCJ50 NATER Ug/E	NYJL-RIMB MBCJ51 WATER Ug/L	NYJL-TRBX2 N/A N/A Ug/L
luajnus	2710	8750	9840	11200		3			MR
·ntiaony	158 E	17.7							Ma
rsenic	452	44.1	33	10.5					HR
arivo	83.1	66.7	107	95.2					MR
eryllium	; ;	į	J	ð					MR
adnive	17.4	. 2	1.4	1.7				31.6	RA
alcive	4770	14700	J	4580					XR
hronium	13.3	19.3	24.2	23.6					HR
obal:	; ;	21.4	3	16					MR
opper	; 1030 E	172 E	317 E				J		ĦR
roh	48700	14700	60200	17200	J	J	125	J	MR
ead	14200	324	58	240					MA
agnesium .	l J	8750	1230	2130					RR
anganese	154	403	340	760					Ħ
ercary	0.9 €	1.1 €		0.36 €					RR
ickel	, ,	29.5	7.8	25.4				J	RR
olassium	1	J	1	J.					NA.
eleniva	13.6 E								MA.
ilver .	125	2.6	•						RR
evibe									PR
halliub	9	R		R					MR
anadium	39.5 E	32.4 E	39.2 [•	,	,	MR
inc _/ anide	847 E	. 166 E	52.6 E	179 €	J	J	1	1	AR NR

ITES:

lank space - compound analyzed for but

- not detected
- estimated value
- estimated value, compound present below CRDL but above IDL
- analysis did not pass EPA QA'QC
- 1 analysis not required

103554

9.49.

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

				(55)	,					
/OLATILES Sample 10 Mm. Traffic Meport Mm. Hatrix Units Dilution Factor Percent Moisture	NYJL-SMI DDK63 MATER Wg/L 1	NYJL-SH2(HS/MSD) BDK64 MATER Wg/L I	NYJL-SM3(DUP) BDR65 MATER Wg/L 1	NYJL-SEDI (MS/MSD) BDR66 SEDIMENT Ug/kg 1 44	NYJL-SED2(DUP) ODR67 SEDIMENT ug/kg i 45	NYJL-RINI BDX60 WATER Wg/L I	NYJL-RINZ BDK69 WATER Wg/L J	NYJL-RINS BDK72 WATER Wg/L I	NYJL-RIN4 DDX73 NATER Vg/L 1	NYJL-FOLK BDPO3 WATER Ug/L I
hToromethane romomethane inyl Chloride hloromethane ethylene Chloride cetone arbon Diswlfide "1-Dichloromethane rans-1,2-Dichloromethane rans-1,2-Dichloromethane				J						
hloroform ,2-Dichloroethane -Butanone ,i,1-Trichloroethane arbon Tetrachloride inyl Acetate rooodichloroethane		•			R					
2-Dichleropropane is-1,3-Dichleropropene ichloroethene ibroeochloronethane 1,2-Trichloroethane		7								•
rans-1,3-Dichloropropene ronofora -Methyl-2-Pentanone -Mexanone strachloroethene oluene ,1,2,2-Tetrachloroethana							•			
ri,c.2-letrachtoroethanu htysbenzene tyrene ylenes (lotal)	1									

OTES:

luve: v2-9005-us

SAMPLING DATE: 5/15/90

PA CASE NO.: 14115 LAB: NET MID-ATLANTIC

lank space - compound analyzed for but not detected

- compound found in lab blank as well as sample, indicates possible/probable blank contamination
- estimated value
- estimated value, compound present below CROL but above IDL
- analysis did not pass EPA 0A/0C
- Presumptive evidence of the presence of the material
- R analysis not required
 - in limite playated if Dilution

1001: 02-9003-01 SAMPLING DATE: 5/15/90

:xachlorobenzene

PA CASE NO.: 14115 LAB: MET MID-ATLANTIC

INULL A SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

SEMI-VOLATILES Sample ID No. INVJE-SWE NYJL-SH2(NS/NSD) NYJL-SH3(OUP) NYJL-SED1(NS/NSD) NYJL-SED2(OUP) NYJL-RIN1 NYJL-RIN2 NYJL-RIN3 NYJL-RIN4 NYJL-RIN4 NYJL-RIN5 BOR65 BORGE 10867 fraffic Report No. **8DE63** BDK64 80148 BDX49 BDE72 80K73 RDP03 tatrix WATER WATER MATER SEDIMENT SEDIMENT WATER MATER MATER WATER WATER wg/L ug/kg ug/kg Inits ug/L wg/L ug/L wg/L ug/L ug/L ug/L ilution Factor/GPC Cleanup (Y) N/A 1 ı 1 1 1 ı 'ercent Moisture 44 45 N/A 'heno ì M sisf2-Chloroethyllether ## ĦŘ 1-Chlorophenol .. 3-Dichlarobenzene .4-Dichlorobenzene enzyl alcohol .2-Dichlorobenzene -Nelaylphenol is(2-Chloroisopropyl)ether -Methylphenal -Mitroso-di-m-dipropylamine exachleroethane itrobenzene sepherene -Hitrophenol .4-Disethylphenol enzoic acid is(2-Chloroethomy)aethane .4-Dichlorophenol ,2,4-Trichlorobenzena aphthalene -Chloroaniline exachlorobuladiene -Chioro-3-Methylphenol ·Methylnaphthalene exachlorocyclopentadiene .4.6-Trichlorophenol ,4,5-frichlorophenol ·Chloronaphthalene ·Mitroaniline isethylphthalate :enaphthylene .4-Dinitrotoluene -Mitroaniline enaphthene .4-Dimitrophenol Hitrophenol ibenzofuran 4-Dinitrotolvene iethylphthalate ·Chlorophenyl-phenyl ether luorene Mitroaniline .4-Dinitro-2-aethylphenol nitrosodiphenylamine Bronophenyl-phenyl ether

Ú

SITE NAME: LI TUMGSTEN 1004: 02-9003-01 SAMPLING DATE: 5/15/90

SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

IPA CASE NO.: 14115 LAB: NET HID-ATLANTIC

GEMI-VOLATILES Gample ID Mo. Traffic Report Mo. Matrix Juits Filution Factor/GPC Cleanup (Y) Percent Moisture	NYJL-SNL BDK63 HATER Ug/L L	NYJL-SH2(MS/MSD) BDK64 WATER Wg/L I	NYJL-SH3(DUP) N BDK65 HATER Ug/L L	YJL-SEDI(MS/MSD) BDR66 SEDIMENT Wg/kg I 44	MYJL-SEDZIDUP) BDK67 SEDIHENT ug/kg 1 45	NYJL-RINI BDK68 WATER Ug/L 1	NYJL-RINZ ODK69 WATER Ug/L 1	NYJL-RIMS BDK72 WATER Ug/L 1	NYJL-RIN4 BDR73 WATER Wg/L I	NYJL-TOLKS BDPO3 MATER Wg/L M/A M/A
'entachlorophenol 'henanthrene inthracene ii-n-butylphthalate luoranthene 'yrene utylbenzylohthalate -,3'-pichloropenzidine enzolalanthracene hrysene isi2-Ethylhexyllohthalate i-n-octylohthalate				J	1	•••				NP. NR NR NR NR NR NR NR NR
enzo(b)fluoranthene enzo(h)fluoranthene enzo(a)pyrene ndeno(1,2,3-cd)pyrene ibenz(a,h)anthracene enzo(g,h,i)perylene										ITR INR INR INR INR INR

OTES:

lank space - compound analyzed for but not detected

- compound found in lab blank as well as sample, indicates possible/crobable blank contamination
- estimated value
- estimated value, compound present below CROL but above IDL
- analysis did not pass EPA QA/QC
- Presumptive evidence of the presence of the material
- ? analysis not required

stection limits elevated if Dilution sctor >1 and/or percent moisture 'Ot

103557

125.6 35.447

Rev. No. 0

TABLE 1
SUMMARY OF SITE INSPECTION ANALYTICAL DATA
(cont'd)

PESTICIDES	:									
Sample ID Mc.	W.JL-SWI	NYJE-SUZENS/MSG1		HYDL-SEDIENS MSDI		NYJL-RINI		NYJL-FIM3		
Traffic Report Hc.	BCK63	BDK64	80865	BDX66	BDK67	96168	8DR69	BDK72	B0x73	BDF03
Matri.	WATER	MATER	VATER	SEDIMENT	SEDIMENT	WASER	WATER	MATER	HATEP	WATEP
Units	1/gr	ug/L	ug/L	ug/tş	wg/kg	ug/L	ug/l	va/l	nė/f	ug/L
Dilution Factor/SPC Cleanup (Y) Percent Moisture		1	1	1 44	1 45					N/A N/A
aloha-BH(• ; • • • • • • • • • • • • • • • • • •	• • • • • • • • • • • • • • • • • • • •		• • • • • • • • • • • • • • • •	••••••		•••	• • • • • • • • • •		KR
beta-ENC	1									MR
delta-BHC										MA
sanna-BHC (Lindane)	i									38
lentach]er										MR
Aldrin										MR.
leotachlor epoxide		•								MR
Endosulfan !	i									MR
ieldrin	i									MR
1.4°-00E	1									HP.
indr in	i	••								NR
indosulfan []		•								MR
1,41-000	•									88
ndosulfan sylfate'	i									##
.4*-901	i									M
lethoxychlor	İ									HR
ndrin ketone	i									WR
icha-Chiordane	•									M
ama-Chlordane	i			•						WR
oxaphene	i			•						MR
roclor-1016	i									RR
roclor-1221	•									MR
roctor-1232	i									MR
roctor-1242	•									MA
roclor-1248	•									MR
roclor-1254	•									MR
roctor-1260	i									NR
OTES:										
lank space - compound analyzed for but										
not detected										
- compound found in lab blank as well as	}									
sample, indicates possible/probable										

 estimated value, compound present below CRQL but above IDL
 analysis did not pass EPA GA/GC
 Presumptive evidence of the presence

blank contamination • estimated value

of the material R - analysis not required etection limits elevated if Dilution actor >1 and/or percent moisture >0%

1004: 01-100: 01

SAMPLING PATE: 5/15/90

EPA CASE NO. 14115 LAB: NET MID-ATLANTIC

TABLE 1 SUMMARY OF SITE INSPECTION ANALYTICAL DATA (cont'd)

INORGANICS Sample ID No. Traffic Report No. Matrix Units	NYJL-SWI MBCP89 WATER Ug/L	NYJL-SN2(HS/NSD) MBCP90 NATER Ug/L	NYJL-SM3(DUP) MBCP91 MATER Wg/L	NYJL-SEDI(HS/MSD) MBCP92 SEDIMENT mg/kg	NYJL-SED2(DUP) MBCP93 SEDINENT mg/kg	NYJL-RINI MBCP94 WATER Wg/L	NYJL-RINZ MBCP95 WATER Wg/L	NYJL-RINS MBCP96 WATER Ug/L	NYJL-RIN4 MBCJ65 WATER Ug/L	RYJL-TOLRI R/A R/A vg/L
Aluminum	j	j	j	1350	610					MR
Antimony		,			R					MR
Arsenic		11	3	72.1	37					NR
3arium	1	3	J	J	J	J	3	1	1	MA
Beryllium				J	J					MR
Cadeiun	•	•		. 2.1						MR
Calcium	90700	53100	74100	202000	308000					WR
Chronium	;			35.4 E	21.5 E					HR
obalt	; 51	366	53	3970	1530					ĦR
lopper	; 25	504	25	610	293					MR
. FOR	; ;	1760	j	2780	1780	J	J			MR
.ead	4.2 €	15.8 E	3.9 €	341	243					MR
lagnesium	1 1	12500	J	3	3					椒
langanese '	1	1649	3	1070	491		3			110
lercury	į			0.23	0.21					KR
ickel	1 1	544	, .	1110	450					MA .
otassi ve	24900 E	22200 E	24700 E	J		3				M
elonius .	į	3								MA.
ilver	İ			14.3	7.6					MR
odive	919000	53100	867000	1720 £	12600 E	J			J	HR
hallium				R	R					KR
anadium	1		j	112 E	57.2 E					XR
inc	30.7	265 E	35.8	782 E	303 E		•			KR
yanide	1	11.7	11.4							MR

OTES:

lank space - compound analyzed for but

- not detected
- estimated value
- estimated value, compound present below CRDL but above IDL
- analysis did not pass EPA QA/QC
- R analysis not required

103559

ev. No. 0

The Upper Glacial Aquifer rests unconformably upon the Raritan Clay of Cretaceous Age. In the Basisan Clay of Cretaceous Age. In the Basisan Clay of Cretaceous Age. In the Basisan Clay of Cretaceous Age. In the Basisan Clay of Cretaceous Age. In the Basisan Clay of Cretaceous Age. In the Basis and the Basis a the vicinity of the site, the Raritan Clay occurs at an approximate depth of 175 feet and has an average thickness of 50 feet. The Raritan Clay occurs at an approximate depth of 1/2 feet and 1/03 and shallow class of 50 feet. The Raritan Clay Consists predominantly of light to dark grey, red, and sing siles and siles white, or yellow clay with variable amounts of silt and fine silty sand. Due to the heterogeneity of sediments within this clay, the permeability is variable; the average vertical permeability is 3.0 x 10.7 cm/sec. Some public supply wells and other private supply wells obtain water from the sandy horizons of the Raritan Clay.

Below the Raritan Clay lies the Lloyd Sand Aquifer of early Cretaceous Age. It can be found at a depth of approximately 225 feet below the surface in the vicinity of the site and is approximately <<>> Teet below the surface in the vicinity of the Site and is approximately 200 feet thick. The Lloyd Sand consists of discontinuous layers of silt, clay, sandy and and analysis and approximately that available in manual life. The average normal hills is 1.7 v. clay, sand, and gravel that exhibit variations in permeability. The average permeability is 1.7 x 10°2 cm/sec. Groundwater within this aquifer flows from north to south. The Lloyd Sand Aquifer serves six public water supply wells within a 3-mile radius of the Li Tungsten Site and has been designated as a sole source aquifer by the U. S. EPA. Ref. Nos. 4 (Volume 1, part 4, pp. 14 to 17) 10, 12, 14, 24, 39

Is a designated sole source aquifer within 3 miles of the site? 3.

A sole source aquifer has been designated within 3 miles of the site. Ref. Nos. 10, 12, 24

What is the depth from the lowest point of waste disposal/storage to the highest seasonal level of the saturated zone of the aquifer of concern?

Contaminants attributable to the facility were detected in samples GW-9 and GW-10, which Were collected by NUS Corporation Region 2 FIT from monitoring well EMW-4, located south of the Mud Pond. The depth to the top of the screened interval in this well has been reported to be 8.6 feet below ground surface. The depth to groundwater below ground surface at the to be o.b teet below ground surface. The depth to groundwater below ground surface at the time of sampling was 5 feet. Therefore, the waste is considered to be in contact with the Ref. Nos. 4, 13, 50

What is the permeability value of the least permeable continuous intervening stratum

The permeability value of the least permeable continuous intervening stratum between the ne permeability value or the least permeable continuous intervening stratum between stratum between the Upper Glacial Aquifer is estimated to be greater than 10-3 cm/sec.

at is the net precipitation for the area?

estimated net precipitation for this area, based upon the normal annual total ipitation minus the mean annual lake evaporation, is approximately 16 inches.

25.6 39.48

7. Identify uses of groundwater within 3 miles of the site (i.e., private drinking source, municipal source, commercial, industrial, irrigation, unusable).

Groundwater within 3 miles of the site is used for private drinking sources, public supply wells, and commercial, industrial, and irrigation applications. Many wells have been closed or have restricted use due to volatile organic chemical contamination from undetermined sources.

Ref. Nos. 9, 12, 15, 38, 39

8. What is the distance to and depth of the nearest well that is currently used for drinking or irrigation purposes?

The nearest well supplying potable water from the aquifer of concern is located 1.3 miles west of the Warehouse. This well (No. 901) is 68 feet deep and is screened within the Upper Glacial Aquifer. Refer to Table 2 for a list of wells within 3 miles of the site.

Ref. Nos. 6, 12, 39

9. Identify the population served by the aquifer of concern within a 3-mile radius of the site.

The population served by the aquifer of concern within a 3-mile radius of the site is approximately 18,000 or more.

Ref. Nos. 9, 12, 16

SURFACE WATER ROUTE

10. Describe the likelihood of a release of contaminant(s) to surface water as follows: observed, alleged, potential, or none. Identify the contaminant(s) detected or suspected, and provide a rationale for attributing the contaminants to the facility.

There is a potential for contaminants to be released to Glen Cove Creek via storm drains on Herb Hill Road and the storm drains on site. Runoff from the landfill and from the main part of the facility enters Glen Cove Creek via these routes. The leaking Mud Holes, Mud Pond, and aboveground 500,000-gallon oil tank are located across the street (Garvies Point Road) from Glen Cove Creek. Chemicals identified in surface soil and groundwater samples around these waste sources include arsenic, selenium, silver, barium, cobalt, chromium, copper, iron, manganese, nickel, strontium, vanadium, zinc, lead, antimony, thallium, aluminum, tungsten, cadmium, titanium, and molybdenum. The site is located on the 100/500-year floodplain; therefore, the potential also exists for surficial contaminants to be transported off site and into Glen Cove Creek via flooding. Surface water samples analyzed for tungsten by inductively coupled plasma mass spectrometry (ICP/MS) were found to contain tungsten, copper, zinc, arsenic, molybdenum, antimony, lead, bismuth, thorium, and uranium.

Ref. Nos. 1, 4 (Volume 1, part 1, pp. 1-14 to 1-15), 8, 21, 22, 27, 28, 31, 32, 37, 40, 43

11. Identify and locate the nearest downslope surface water. If possible, include a description of possible surface drainage patterns from the site.

The nearest downslope surface water is Glen Cove Creek, which generally flows southwest but is also affected by the tides. Glen Cove Creek is adjacent to the south property boundary; it then flows into Hempstead Harbor and Long Island Sound. Runoff from the site and from storm drains on Herb Hill Road can drain directly into Glen Cove Creek via several outfalls.

Ref. Nos. 6, 8, 27, 43, 44

WELLS WITHIN 3 MILES OF LI TUNGSTEN, GLEN COVE, NY

UN = Unused, P.S. = Public Supply, IND = Industrial, COM = Commercial, IRR = Irrigation, UNK = Unknown, AC = Air Conditioning

**:	_	commercial, IRR = Irrigat
Well No.		····yat
100	Use	_
109		<u>Contaminated</u>
110	UN	
112 114	UN	
115	UN	
116	IRR	
117	UN	
	UN	
119	UN	
120	P.S.	_
121	UN	•
121A	UN	
6 60	IRR	
661	IND	
801-818	UN	
834	UN	
835	UN	
842	P.S.	
901	UN	
902	P.S.	
903	P.S.	
904	P.S.	
905-909	UN	
1037	P.S.	
1149-1153	P.S.	
1171-1174	UN	
1327	UN	
1595	P.S.	
1651	P.S.	
1917	P.S.	•
2027	IND	
2060	UN	
2087	UN	
2316	IND/UN	
2616	IND	
3310	IRR	×
3466	IND	
3892	P.S.	
4432	P.S.	×
4440	COM	â
4462	DOM	^
4639	UN	•
5071	UNK	
5201	IRR	
5250	P.S.	
5261	UN	
- •	P.S.	
	· .J.	V
		×

TABLE 2 (CONTINUED)

Ref. 6

Well No.	<u>Use</u>	Contaminated
5450	IRR	
5762	P.S.	
5792	P.S.	
6289	UN	
6289	IRR	
6416	UN	
6444	IRR	
6549	IND	
6579	UNK	X
6587	UN	
6665	UN	
6668-70	UN	•
6708	UN	
6806	IRR	
6881	UN	
6883	UN	
6973	UN	v
7427 7430	IND	X
7439 7614	UNK IND	
7664	IRR	×
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7834	IRR	
7857	P.S.	
8048	UNK	
8224	IND	
8259	UN	
8326	P.S.	X
8327	P.S.	×
8394	UNK	
8690	UNK	
8709	IND	
8716	UN	
8887	IND	×
8898	UN	
8937	COM	
9066	UN	
9100	UN	
9115	UN	
9117	UN	
9210 :	P.S.	
9211	P.S.	
9334	P.S.	

Ref. Nos. 10, 12, 39

12. What is the facility slope in percent? (Facility slope is measured from the highest point of deposited hazardous waste to the most downhill point of the waste area or to where contamination is detected.)

The facility slope, as measured from the northern boundary of the landfill to the southern boundary of the landfill, is 5 percent.

Ref. Nos. 4, 6,13

13. What is the slope of the intervening terrain in percent? (Intervening terrain slope is measured from the most downhill point of the waste area to the probable point of entry to surface water.)

The slope of the intervening terrain, as measured from the southern boundary of the landfill to Glen Cove Creek, is less than 2 percent.

Ref. Nos. 6, 8, 13

14. What is the 1-year 24-hour rainfall?

The 1-year 24-hour rainfall for the area is approximately 3 inches.

Ref. No. 14

15. What is the distance to the nearest downslope surface water? Measure the distance along a course that runoff can be expected to follow.

Glen Cove Creek is adjacent to the southern property boundary. Previously permitted outfalls and on-site storm drains discharge through the bulkhead along the southern property boundary directly into Glen Cove Creek.

Ref. Nos. 8, 13, 27, 43, 44

16. Identify uses of surface waters within 3 miles downstream of the site (i.e., drinking, irrigation, recreation, commercial, industrial, not used).

Surface water uses within 3 miles downstream of the site include recreational and commercial.

Ref. Nos. 6, 20

17. Describe any wetlands, greater than 5 acres in area, within 2 miles downstream of the site. Include whether it is a freshwater or coastal wetland.

No wetlands greater than 5 acres in area have been identified within 2 miles downstream of the site.

Ref. Nos. 6, 20

18. Describe any critical habitats of federally listed endangered species within 2 miles of the site along the migration path.

No critical habitats of federally listed endangered species have been identified within 2 miles of the site. However, Hempstead Harbor is a waterfowl wintering area most noted for scaup, canvasback, and black ducks, and is a nursery/feeding habitat for striped bass, bluefish, Atlantic

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silverside, menhaden, winter flounder, and blackfish. Hempstead Harbor has been designated as a "significant coastal fish and wildlife habitat" by the NYS Department of State under Policy 7 of the Waterfront Revitalization and Coastal Resources Act of 1981.

Ref. Nos. 6, 7, 20, 23, 25

19. What is the distance to the nearest sensitive environment along or contiguous to the migration path (if any exist within 2 miles)?

No sensitive environments have been identified along Glen Cove Creek or Hempstead Harbor within 2 miles of the site.

Ref. Nos. 6, 7, 20, 23, 25

20. Identify the population served or acres of food crops irrigated by surface water intakes within 3 miles downstream of the site and the distance to the intake(s).

There are no crops irrigated by surface water intakes within 3 miles downstream of the site.

Ref. Nos. 6,11

21. What is the state water quality classification of the water body of concern?

The state water quality classification for Hempstead Harbor north of Bar Beach is Class SA (suitable for shellfishing for market purposes and primary/secondary recreation). The state water quality classification for Glen Cove Creek is Class 1 (secondary contact recreation except for primary recreation and shellfishing).

Ref. No. 18

22. Describe any apparent biota contamination that is attributable to the site.

Biota contamination attributable to the site exists along the grassy areas around the Mud Pond, Mud Holes, and the nine waste piles. There was a notable lack of vegetation around these areas, and grass near the fence along Garvies Point Road was stained black.

Ref. Nos. 4, 5, 13

AIR ROUTE

23. Describe the likelihood of a release of contaminant(s) to the air as follows: observed, alleged, potential, none. Identify the contaminant(s) detected or suspected, and provide a rationale for attributing the contaminant(s) to the facility.

There is a potential for release of contaminants from the site into the air. Tank covers, siding shingles, roofing tiles, and pipe wrapping, all of which are known to contain asbestos, are in a state of decay. However, analyses of indoor and outdoor air samples previously collected from the site indicate the presence of little or no volatile organic chemicals and airborne metals; analyses also indicate little or no asbestos particulates. Larger pressurized tanks containing aqueous ammonia and propane, and open-air tanks containing hydrochloric acid and tungsten acid could potentially release their contents to the air.

Ref. Nos. 4 (Volume 1, part 1, pp.1-6; Volume 2, part 6), 13, 25, 26, 34, 35, 36, 41

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24. What is the population within a 4-mile radius of the site?

The population within a 4-mile radius of the site is approximately 67,900.

Ref. No. 17

FIRE AND EXPLOSION

25. Describe the potential for a fire or explosion to occur with respect to the hazardous substance(s) known or suspected to be present on site. Identify the hazardous substance(s) and the method of storage or containment associated with each.

There is a potential for fire or explosion to occur with respect to the hazardous substances known to be present on the site. An outdoor, partially filled and pressurized tank of propane gas and a pressurized tank of aqueous ammonia are present on site and represent a potential for an explosion or fire to occur. In a letter to the NYSDEC dated January 31, 1990, the mayor of the City of Glen Cove expressed his concern about the potential for a fire to occur at the site and for the safety of local fire fighters who would have to enter the site, if such an event were to occur.

Ref. Nos. 4 (Part 6), 13, 25, 26, 33, 41

26. What is the population within a 2-mile radius of the hazardous substance(s) at the facility?

The population within a 2-mile radius of the hazardous substances present on the site is approximately 35,400.

Ref. No. 17

DIRECT CONTACT/ON-SITE EXPOSURE

27. Describe the potential for direct contact with hazardous substance(s) stored in any of the waste units on site or deposited in on-site soils. Identify the hazardous substance(s) and the accessibility of the waste unit.

There is a potential for direct contact with the hazardous substances deposited in on-site soils, which include heavy metals, PCBs, and radioactive elements. Along Garvies Point Road, the Mud Pond has overflowed and stained the soil. The stained soil contains notable concentrations of arsenic, antimony, chromium, copper, lead, mercury, vanadium, and zinc.

Ref. Nos. 4, 5, 13, 21, 33, 36, 49, 50

28. How many residents live on a property whose boundaries encompass any part of an area contaminated by the site?

There are no residents who live on a property whose boundaries encompass any part of an area contaminated by the site.

Ref. Nos. 5, 13, 50

29. What is the population within a 1-mile radius of the site?

The population within a 1-mile radius of the site is approximately 9,900.

Ref. No. 17

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PART V: ACTUAL HAZARDOUS CONDITIONS

Waste processed ore containing heavy metals and radioactive isotopes of uranium, thorium, and radium occur in nine waste piles and within many drums and crates in many parts of the site. The drums and crates are badly weathered, corroded, and spilling their contents on the ground. Particulates from the waste piles can become airborne as they are not covered. The waste piles also contain radioactive slags mixed with the soil, and several separate piles containing large chunks of radioactive slag have been roped off and marked with radiation placards; however, radioactive slag is not considered a hazardous waste in New York State. The Li Tungsten Site has been designated as a Class 2 site (significant threat to public health or environment) on the NYSDEC registry because of the presence of other contaminants. The stained soil on the corner of Garvies Point Road contains notable concentrations of many heavy metals. Several 55-gallon drums containing elevated levels of radioactive waste process ore and soil that was excavated from behind the wooden fence at the corner of Garvies Point and Herb Hill Roads have been stored within the Dice Building until an approved disposal site for New York State has been established. Tank covers, pipe wrappings, wallboard, and shingles on site contain asbestos. These items are known to be in poor condition and have been found in broken pieces upon the ground. Although air testing previously conducted has not shown the presence of airborne asbestos particles, the potential exists for a release of particulates to the air. Also, many wooden, steel, or fiberglass tanks still contain process solutions containing heavy metals and concentrated or spent acids and bases. There are two pressurized tanks on site, one of which contains aqueous ammonia and the other propane gas. Although the site is patrolled by a one-man private security force, the site is very large and the fence surrounding the site has been broken many times; therefore, there is a potential for unauthorized entry to the site. The buildings on the site are in poor condition and local officials have expressed a concern for the safety of their firemen; they are especially concerned about the asbestos dusts and particulates that may be released if a fire should break out on the site. At least four different contaminant plumes have been identified as a result of several groundwater sampling events conducted on site.

No other actual hazardous conditions pertaining to human or environmental contamination have been documented. Specifically:

- Contamination has not been documented either in organisms in a food chain leading to humans or in organisms directly consumed by humans.
- There have been no documented observed incidents of direct physical contact with hazardous substances at the site involving a human being (not including occupational exposure) or a domestic animal.

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• There have been no documented incidents of damage to fauna (e.g., fish kill) that can be attributed to the hazardous materials at the facility.

Ref. Nos. 4, 13, 25, 26, 33, 36, 49, 52

PART VI: SITE SUMMARY AND RECOMMENDATIONS

Li Tungsten is located in an industrial area on approximately 26 acres along the north bank of Glen Cove Creek in the City of Glen Cove, Nassau County, New York. From the 1940s to the early 1980s, tungsten ores imported from Mainland China and Canada were smelted at this facility for the making of tungsten carbide powder, tungsten wire, and welding rods. In 1985, the company filed for bankruptcy; the property is presently owned by the Glen Cove Development Company located in Baltimore, Maryland.

Although the site is presently inactive, most of the wastes generated by the facility remain on site. These wastes include 17,000 tons of solid residue/ore materials in piles, in a landfill, in wooden crates, and in 30- and 55-gallon drums. Some of the drums are overstacked and some have toppled and have broken open, spilling their contents upon the ground. One hundred and eight drums containing acids, waste oil, and organics have been overpacked and/or staged to a secure area on site. The remaining unsound drums are also recommended for overpacking to eliminate the potential for a release of their contents. Elsewhere on the site, there are approximately 373,000 gallons of various liquids stored in 224 aboveground tanks of unknown physical condition, some of which contain hazardous organic and inorganic liquids. The inorganic liquids include spent or unused hydrochloric acid and aqueous ammonia. Fifty tanks have been inspected for leaks and rupture. Two tanks were determined not to be secure and have been drained and their contents drummed for disposal. Small quantities of identifiable chemicals have been overpacked and secured, while small quantities of unidentified chemicals remain in some areas. Thirty-eight electrical transformers formerly located on site, three of which contained PCB-contaminated oil, have been drained, drummed, and disposed of at a licensed off-site facility. Removal activities have also begun with respect to some of the surficial containers (including pressurized cylinders).

A site investigation conducted by a consulting firm on behalf of the site owner was completed in May of 1988, during which samples were taken from 10 existing groundwater monitoring wells and 13 more monitoring wells were installed. Analyses of samples from these wells identified four underground plumes within the groundwater of the Upper Glacial Aquifer. One plume occurs at a depth of approximately 20 feet along the eastern boundary of the site and was found to contain several dry cleaning solvents related to tetrachloroethylene. The plume is believed to originate from a dry cleaning facility that formerly occupied the property adjacent to the site. Another plume was found along the western boundary of the site and was traced to an adjacent property formerly occupied by a petrochemical company. Both plumes are moving south towards Glen Cove Creek. Another plume of No. 2 fuel oil occurs in the vicinity of a leaking 500,000-gallon tank north of Garvies Point Road. The last plume is located around the Mud Pond/Mud Holes, which contain waste processing water and heavy metals. Chloride and sulfate compounds, and notable concentrations of

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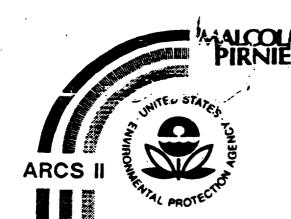
PART VI: SITE SUMMARY AND RECOMMENDATIONS (CONT'D)

lead, cadmium, tungsten, chromium, arsenic, barium, and silver have been detected in groundwater samples collected from this area. The materials leaking from the fuel oil tank and the ponds have also scarred the vegetation and stained the soil in this area. Asbestos fibers from decaying tank covers and pipe wrapping materials are known to be present on the ground. Similarly, waste piles containing raw and processed tungsten ores are known to contain radioactive radium, uranium, and thorium compounds used in the ore refining process. The United States Environmental Protection Agency issued an Administrative Order on Consent to the Glen Cove Development Company on July 21, 1989, outlining initial actions to be taken at the site. The site is scheduled for a cleanup of hazardous wastes including, but not limited to, the removal of drums, the contents of the tanks, and the laboratory chemicals, but plans for cleanup of the groundwater and soil have not been finalized. Development as a residential area is planned for the site.

Analytical results from groundwater, surface water, soil, and sediment samples collected from this site by NUS Corporation Region 2 FIT in April and May of 1990 indicate a release of significant concentrations of contaminants associated with tungsten refining to the environment. Elevated concentrations of antimony, arsenic, barium, beryllium, copper, cobalt, chromium, lead, manganese, mercury, nickel, vanadium, cadmium, uranium, thorium, molybdenum, bismuth, zinc, and cyanide were detected in soil and/or groundwater samples. The uranium and thorium compounds are known to be radioactive. Analytical results from the surface water and sediment samples collected from onsite waste sources indicate the presence of notable concentrations of PCBs (Aroclor-1248 and Aroclor-1254) and elevated concentrations of metals. Two of the surface water samples collected from on-site waste sources contained cyanide. Surface water samples collected from Glen Cove Creek showed the presence of tetrachloroethene; sediment samples collected from the creek contain polycyclic aromatic hydrocarbons and elevated concentrations of several metals.

Based upon the high target population potentially affected by groundwater contamination and the potential for direct contact with some of the wastes on site, the Li Tungsten Site is recommended for a LISTING SITE INSPECTION. All of the radioactive waste piles should be roped off from unauthorized access and labelled with radiation placards until they can be contained/covered, removed from the site, and properly stored at a licensed facility. The propane and aqueous ammonia tanks and other large process tanks containing organic and inorganic liquids/residues should be emptied and disposed of properly. The remaining drums, barrels, and crates of tungsten ore/residues should be recycled or processed at another tungsten refining facility. The many empty 55- and 30-gallon drums on site should be crushed and properly disposed. Lastly, a cleanup plan for the contaminated groundwater, surface water, soil, and sediment should be formulated.

REFERENCE 7



Contract No. 68-W9-0051

LI TUNGSTEN GLEN COVE, NEW YORK

Work Assignment No. 025-2L4L

REMEDIAL INVESTIGATION/FEASIBILITY STUDY WORK PLAN PART I OF II

Remedial Planning Activities at Selected Uncontrolled Hazardous Substance Disposal Sites USEPA Region II (NY, NJ, PR, VI)

Malcolm Pirnie, Inc.

2 Corporate Park Drive
White Plains, New York 10602

March 1993

WORK PLAN REMEDIAL INVESTIGATION/FEASIBILITY STUDY

PART I OF II

LI TUNGSTEN
GLEN COVE, NEW YORK

MARCH 1993

ARCS Contract No. 68-W9-0051
USEPA Work Assignment No. 025-2L4L

MALCOLM PIRNIE, INC. 2 Corporate Park Drive White Plains, New York 10602

ARCS II CONTRACT NO. 68-W9-0051

WORK ASSIGNMENT # 025-2L4L

SITE NAME:

LI LUNGSTEN

RI/FS WORK PLAN

MARCH 1993

CONTRACTOR QA/QC SIGN-OFF

Malcolm Pirnie, Inc. has reviewed this draft document in accordance with the contractor's ARCS II QAPP and is submitting it to USEPA, Region II in compliance with the requirements under Work Assignment No. 025-2L4L and Contract No. 68-W9-0051.

This document has not been approved by USEPA Region II and is not intended for release to the public.

Dennis G. McGrath

SITE MANAGER

Date: 18 Nach 1993

S.K. Krishnaswami

ARCS II PMO PROGRAM MANAGER

Date: March 19,

LI TUNGSTEN RI/FS WORK PLAN

EXECUTIVE SUMMARY

SITE LOCATION AND BACKGROUND

The Li Tungsten site is located at 63 Herb Hill Road in the City of Glen Cove, Nassau County, Long Island, New York. This site has a complex history of name and ownership changes, and environmental site assessments, investigations and removal actions. Specific details are discussed in Section 2.0. From early 1940's until approximately 1985, tungsten ores or concentrates, imported primarily from mainland China, South America and Canada, were smelted at this facility for the production of tungsten carbide powder, tungsten wire, and welding rods (NUS, 1989; 1990). In 1985 the company filed for bankruptcy and the facility ceased operation.

Large quantities of the ore concentrates were left on site in various processed and unprocessed forms. The ore which is present in drums, wooden crates and piles both inside and outside the buildings, contains heavy metals and radioactive isotopes of uranium, thorium, and radium. Many of the drums and crates located outside are weathered and/or corroded to a point where the contents have spilled on the ground. In other areas, the drums have been over-stacked and have become very unstable as the drums deteriorated and corroded.

Numerous aboveground wooden, steel or fiberglass tanks were used during the various smelting processes, and to store reactants (e.g., hydrochloric acid, ammonia, hydrogen) and/or intermediate compounds (e.g., ammonium paratungstate or APT). Some of these tanks may still contain some hazardous and inorganic liquids. As the tungsten ore moved through its various processing stages, the radioactive isotopes became more concentrated in the residual waste or slag. There are indications that some of this slag was placed in waste piles at the ground surface and/or buried on site (NUS, 1989; 1990). Heavy metals which constitute impurities that were removed during the extraction process include: antimony, arsenic, barium, bismuth, copper, cobalt, chromium, lead, manganese, mercury, molybdenum, nickel, thorium, uranium, vanadium, and zinc.

Several of the buildings on site have deteriorated to a point where they represent a physical safety hazard. Portions of some walls and roofs have collapsed. In addition, friable and non-friable asbestos is present as pipe wrap, tank insulation, siding shingles, and roof tiles. Standing water in the West Dice Building has flooded and concealed a deep pit in the floor.

Previous Site Investigations

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Various site investigation activities were conducted at the site between 1988 and 1990 by the Nassau County Department of Health (NCDOH), the New York State Department of Environmental Conservation (NYSDEC), the potentially responsible parties (PRPs), and the United States Environmental Protection Agency (USEPA). Results of these sampling activities have indicated the presence of heavy metals, fuel oil constituents, and volatile organics in the groundwater, surface water, sediments and soils.

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Current Conditions

The Li Tungsten site ceased operations in June 1985 and has been inactive since. Site security (fencing and guard) was addressed as one of the interim remedial measures in the AOC. Although a one person security guard is maintained on a 24-hour basis, the site could be entered without the knowledge of the security force through breaks in the fence. During the site tour, observations were made that vandalism has occurred. Many of the salvageable fixtures (e.g., copper wiring and piping) have been removed and general debris (e.g., washing machines, mattresses) have been left behind.

OBJECTIVE OF THE RI/FS

This Remedial Investigation/Feasibility Study (RI/FS) is designed to collect sufficient data on the nature and extent of contamination to remediate the site. In achieving this objective, these data will be used to determine contamination sources, identify migration pathways, perform an assessment of human health and ecological risks, and support the selection of remedial alternatives to mitigate or reduce risks in accordance with the requirements of the National Contingency Plan (NCP) and the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA), as amended by the Superfund Amendments and Re-authorization Act of 1986 (SARA).

The Health and Safety Plan (HASP), and Field Operations Plan (FOP), which includes the Quality Assurance Project Plan (QAPjP) and the Field Sampling Plan (FSP), will be prepared after the Work Plan has been approved by the USEPA.

INITIAL EVALUATION

The contamination at the Li Tungsten site exists in the groundwater, soil, surface water and sediments. The groundwater contains VOCs and inorganic compounds. The major VOCs contamination is present in two areas and may be related to two off-site sources. The inorganic contamination on-site is a result of the past facility operations and disposal practices. Drums, crates and piles of processed ore and slag will continue to act as contaminant source to the groundwater until they are removed. The disposal area in Parcel B, the two Mud Holes, the Mud Pond and the storm drains are also potential contaminant sources.

The surface water contamination consists mostly of inorganic compounds and relative low levels of VOCs. Continuing sources to surface water contamination consists of runoff from the residual ores, the disposal area in Parcel B, and the storm drains.

During site visits, several safety related observations were made. These observation related to obstructions and site conditions that would affect worker safety in the performance of RI field investigation tasks. To eliminate these safety hazards, we propose that additional interim remedial actions be implemented to address each of the safety hazards, before RI field investigation tasks are initiated.

6 of 27

1.0 INTRODUCTION

1.1 Overview

The Li Tungsten Corporation (Li Tungsten) site is an inactive 26 acre site located at 63 Herb Hill Road, City of Glen Cove, Nassau County, New York (USEPA ID #NYD9868826-60). From early 1940's until approximately 1985, tungsten ores or concentrates, imported primarily from mainland China, South America and Canada, were smelted at this facility for the production of tungsten carbide powder, tungsten wire, and welding rods (NUS, 1989; 1990). In 1985 the company filed for bankruptcy and the facility ceased operation.

Large quantities of the ore concentrates were left on site in various processed and unprocessed forms. The ore which is present in drums, wooden crates and piles both inside and outside the buildings, contains heavy metals and radioactive isotopes of uranium, thorium, and radium. Many of the drums and crates located outside are weathered and corroded to a point where the contents have spilled on the ground. In other areas, the drums have been overstacked and have become very unstable as the drums deteriorated and corroded. Since many of the drums contain radioactive material, they represent both a potential health hazard as well as a physical safety hazard.

The amount of extractable tungsten in a specific ore is dependent on the ore characteristics and the mineral assemblages of the ore. While tungsten occurs in 29 known mineral species, numerous isomorphous substitutions are possible within the tungsten minerals. It was necessary during the smelting, therefore, to be able to vary the extraction process to separate the various accessory metals (or impurities) depending upon the specific type of ore or concentrate that was imported. The smelting was generally conducted in relatively small batches, to permit any individual or combination of extraction treatments. Typical treatments in the smelting included physical, chemical and mechanical processes including sizing and crushing; gravity, magnetic and electrostatic separation; roasting; leaching; floatation; and fusion. An analytical laboratory was located on site to perform chemical analysis on the ore and pilot testing of the extraction treatments.

Numerous aboveground wooden, steel or fiberglass tanks were used to perform the extraction treatments and to store reactants (e.g., hydrochloric acid, ammonia, hydrogen)

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and/or intermediate compounds (e.g., ammonium paratungstate or APT). Some of these tanks may still contain some hazardous and inorganic liquids. As the tungsten ore moved through its various processing stages, the radioactive isotopes became more concentrated in the residual waste or slag. There are indications that some of this slag was placed in waste piles at the ground surface and/or buried on site (NUS, 1989; 1990). Heavy metals which constitute impurities that were removed during the extraction process include: antimony, arsenic, barium, bismuth, copper, cobalt, chromium, lead, manganese, mercury, molybdenum, nickel, thorium, uranium, vanadium, and zinc.

Many of the buildings on site have deteriorated to a point where they are not considered safe to enter. Portions of some walls and roofs have collapsed. Friable and non-friable asbestos is present as pipe wrap, tank insulation, siding shingles, and roof tiles. Standing water in the West Dice Building has flooded a deep pit in the floor.

Various site investigation activities were conducted at the site between 1987 and 1990 by the Nassau County Department of Health (NCDOH), the New York State Department of Environmental Conservation (NYSDEC), the potentially responsible parties (PRPs), and the United States Environmental Protection Agency (USEPA). Results of these sampling activities have indicated the presence of heavy metals, fuel oil constituents, and volatile organics in the groundwater, surface water, sediments and soils.

This Remedial Investigation/Feasibility Study (RI/FS) is designed to collect sufficient data on the nature and extent of contamination to remediate the site. In achieving this objective, these data will be used to determine contamination sources, identify migration pathways, perform an assessment of human health and ecological risks, and support the selection of remedial alternatives to mitigate or reduce risks in accordance with the requirements of the National Contingency Plan (NCP) and the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA), as amended by the Superfund Amendments and Reauthorization Act of 1986 (SARA).

1.2 Approach to Development of Work Plan

Malcolm Pirnie, Inc., (MPI) is submitting this Work Plan to the USEPA in response to Work Assignment #025-2L4L under the Alternative Remedial Contracting Strategy (ARCS)

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Contract No. 68-W9-0051. This Work Plan presents the proposed technical scope of work for the RI/FS and includes a schedule for the performance of the work.

This Work Plan has been prepared in accordance with current USEPA guidance. The following are several of the documents specifically applicable to preparation of an RI/FS that were considered in preparing this Work Plan:

- Interim Final Guidance for Conducting Remedial Investigations and Feasibility Studies Under CERCLA, OSWER Directive 9355.3-01. (USEPA, 1988a)
- Data Quality Objectives: Development Guidance for Uncontrolled Hazardous Waste Site Remedial Response Activities, OSWER Directive 9355.0-7B, (USEPA, 1987a).
- Interim Guidance of Superfund Selection of Remedy, OSWER Directive 9355.0-19, (USEPA, 1986a).
- Additional Interim Guidance for FY-87 Records of Decision, OSWER Directive 9355.0-21, (USEPA, 1987b).
- Risk Assessment Guidance for Superfund, Volume I, Human Health Evaluation Manual Part A (USEPA, 1989a).
- Risk Assessment Guidance for Superfund, Volume II, Environment Evaluation Manual (USEPA, 1989b).
- Superfund Exposure Assessment Manual (USEPA, 1986b).
- Draft Generic Work Plan Guidance (USEPA, 1989c).
- CERCLA Compliance with Other Laws Manual, Interim Final. EPA/540-/G-89/006. Office of Emergency and Remedial Response, Washington, D.C. August 1988, 195 pp, (USEPA, 1988b).
- Guide for Conducting Treatability Studies Under CERCLA (Interim Final)
 EPA/540/2-89/058, December 1989, 138 pp, (USEPA, 1989d).

Preparation of this Work Plan was based upon a review and consideration of data, information, and discussions related to the following:

• Two site visits by MPI personnel on September 1, 1992 and February 3, 1993.

- USEPA comments on the Draft Work Plan, letter dated December 24, 1992 and subsequent discussions.
- Scoping meeting with the USEPA held on September 3, 1992.
- Li Tungsten Site Investigation Report. Prepared for Compon Reality Corporation, New York, NY by RTP Environmental Associates, Inc., Westbury, NY, May 1988, 2 volumes (RTP, 1988).
- Final Draft, Preliminary Assessment, Li Tungsten, Glen Cove, NY. Revision No. 1 dated October 18, 1989 with Appendices (NUS, 1989).
- Final Draft, Site Inspection Report, Li Tungsten, Glen Cove, NY. September 28, 1990 with Appendices (NUS, 1990).
- Interim Remedial Actions Report. Prepared for Glen Cove Development Company, April 4, 1990 (HART, 1990).
- Final Remedial Investigation Report, Mattiace Petrochemical Site, Operable Unit One, Glen Cove, NY. Volumes I and II (EBASCO, 1991).
- Topographic Map Sea Cliff, NY Quadrangle, 1:24,000, Photorevised 1979 (USGS, 1979).

1.3 Scope of Work

The scope of work for this Work Plan was outlined in the Work Assignment Form and Statement of Work which was transmitted to MPI from the USEPA in a letter from the Contracting Officer (CO) dated August 26, 1992. The Statement of Work identified the following tasks:

- Review existing background documents provided by USEPA.
- Develop an RI/FS Work Plan that is comprehensive enough to support a Record of Decision (ROD) for the entire study area.
- Attend scoping meeting within 10 days after issuance of the work assignment.

1.4 Work Plan Content

This Work Plan is organized into nine sections of text including references and a glossary. A brief description of each section follows.

127

Section 1.0, INTRODUCTION, presents an overview of the environmental conditions at the site, the approach used in developing the Work Plan, the scope of work, and the organization and content of the Work Plan.

Section 2.0, SITE BACKGROUND AND SETTING, presents the background of the site including the location, history and current conditions.

Section 3.0, INITIAL EVALUATION, presents an initial evaluation of the existing data base. This section includes a description of the types of waste present, site hydrogeology, climate, population and environmental resources, migration and exposure pathways, a preliminary identification of applicable or relevant and appropriate requirements (ARARs), a preliminary assessment of public health and environmental impacts, a summary of additional data requirements, remedial action objectives, and recommendations for interim remedial actions to be completed before the RI is initiated.

Section 4.0, WORK PLAN RATIONALE, includes the Data Quality Objectives (DQOs) for RI sampling and analytical activities, and the approach for preparing the Work Plan, which illustrates how the activities will satisfy data needs.

Section 5.0, TASK PLANS FOR RI/FS, presents a proposed scope for each standard task of the RI/FS in accordance with the RI/FS guidance document (USEPA 1988a).

Section 6.0, PROJECT SCHEDULE, presents the anticipated schedule for the RI/FS tasks.

Section 7.0, PROJECT MANAGEMENT APPROACH, presents project management considerations that define relationships and responsibilities for selected task and project management teams.

Section 8.0, REFERENCES, provides a list of references used to develop material presented in this Work Plan.

Section 9.0, GLOSSARY OF ABBREVIATIONS, provides a glossary of abbreviations and acronyms used in this Work Plan.

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The Health and Safety Plan (HASP), and Field Operations Plan (FOP), which includes the Quality Assurance Project Plan (QAPjP) and the Field Sampling Plan (FSP), will be prepared after the Work Plan has been approved by the USEPA.

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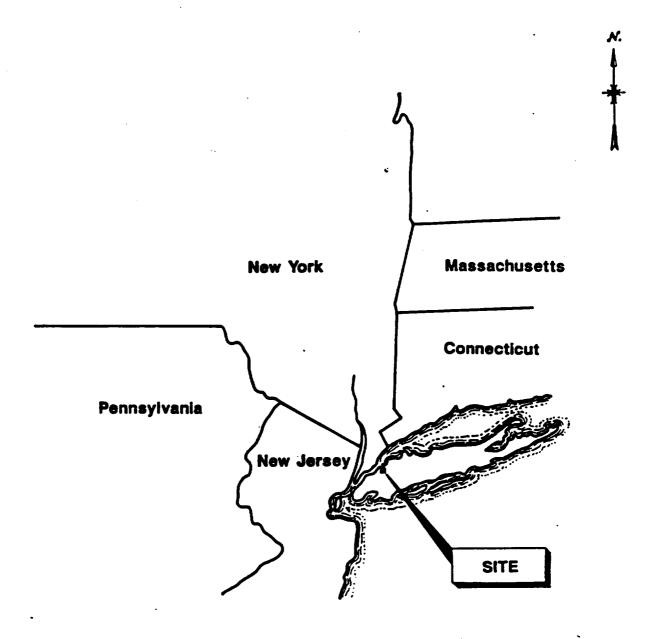
2.0 SITE BACKGROUND AND SETTING

2.1 Site Location

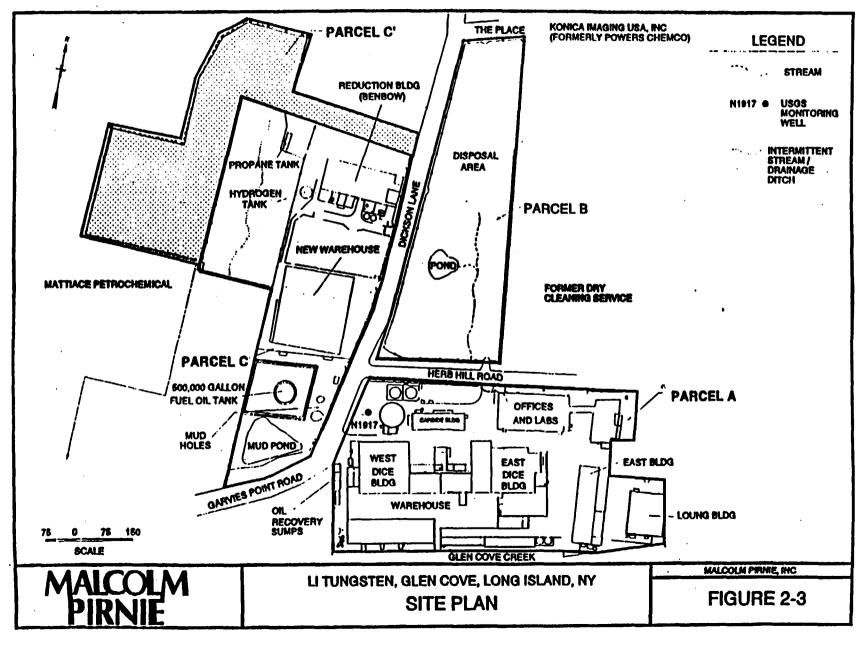
The Li Tungsten site is located at 63 Herb Hill Road in the City of Glen Cove, Nassau County, Long Island, New York. A regional map and a site location map are provided in Figures 2-1 and 2-2, respectively. The geographic coordinates of the site are latitude 40°51'36" North and longitude 73°38'25" West. Also located on Figure 2-2 is the adjacent Mattiace Petrochemical site which is on the National Priorities List (NPL) and was the subject of a recently completed an RI/FS directed by the USEPA (EBASCO, 1991).

The site is approximately 26 acres and consists of four (4) separate parcels designated A, B, C and C'. For the purpose of this Work Plan and subject to the findings of the field investigation, the study area is defined as the entire 26 acres. The location of Parcels A, B, C and C' and the significant site features on each parcel are shown on the site plan in Figure 2-3.

Parcel A is approximately seven acres and served as the main operations center when the site was active. It contains the majority of buildings, structures (e.g., tanks, two surface impoundments) and drums/crates of tungsten ore. It is bounded by Herb Hill Road on the north, Garvies Point Road on the west, an adjoining property on the east, and Glen Cove Creek on the south. Parcel B is the smallest of the three parcels, approximately six acres, and is located due north of Parcel A. Parcel B is bounded by Herb Hill Road on the south, Dickson Lane on the west, The Place on the north, and an adjoining property on the east. The area south of the pond on Parcel B was used primarily as a parking lot when the plant was active, however, disposal activities also are believed to have taken place north of the pond (RTP, 1988). The disposal area north of the pond on Parcel B has been referred to in previous reports (HART, 1990; NUS, 1989, 1990, 1991) as a "landfill". Observations made during the second site visit confirmed that disposal activities have taken place in that portion of Parcel B, but insufficient information is available to confirm that actual landfilling operations took place. Further references to this area in the Work Plan text and on figures, therefore, will refer to it as a disposal area. Parcel C is the largest of the three parcels, approximately 14 acres, however, not all of this parcel was part of the Li Tungsten property during active site operations. The Glen Cove Development Corporation (GCDC) acquired



MALCOLM PIRNIE LI TUNGSTEN
GLEN COVE, NEW YORK
REGIONAL LOCATION MAP



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approximately four acres of undeveloped property, designated Parcel C', sometime after 1984. Parcel C contains several buildings, a 500,000 gallon aboveground fuel oil storage tank, and three surface impoundments (e.g., Mud Pond and two Mud Holes) used to dispose of process waste water.

2.2 Site History

This site has a complex history of name and ownership changes, and environmental site assessments, investigations and removal actions. Specific details are discussed in the paragraphs below. The chronological history of site ownership, operations, and preliminary investigations/interim remedial actions is summarized in Table 2-1.

Early in the 1940's the National Reconditioning Company was formed by Kuo Ching (K.C.) Li. The company was operated and managed by the Wah Chang Trading Corporation of New York. In addition to being the chairman and chief engineer of Wah Chang Trading Corporation, K. C. Li was also a distinguished mining engineer, discoverer of tungsten in China, and was responsible for first importing tungsten into the United States. The purpose of the company was to build a facility in Glen Cove, NY, to concentrate tungsten ores.

The facility became operational in 1942. Operation consisted of processing raw ore and scrap tungsten concentrates to produce ammonium paratungstate (APT) and subsequently formulating APT to metal tungsten powder and tungsten carbide powder. Other specialty products that were produced included: tungsten carbide powder for plasma spraying; tungsten titanium carbide powder; tantalum carbide powder; tungsten spray powder; crystalline tungsten powder; and molybdenum spray powder.

Based on available information, a variety of extraction processes (or treatments) were used to separate the various accessory metals (or impurities) from the tungsten depending upon the specific type of ore or concentrate that was imported. The smelting was generally conducted in relatively small batches, to permit any individual or combination of extraction treatments. Typical treatments in the smelting process included physical, chemical and mechanical processes such as: sizing and crushing; gravity, magnetic and electrostatic

TABLE 2-1 CHRONOLOGY OF EVENTS AND SITE INVESTIGATIONS LI Tungsten Site Glen Cove, New York

DATE	EVENT
1940	National Reconditioning Corporation was formed by K. C. Li with the express purpose of building the Glen Cove facility.
1942	Facility becomes operational. Operation consisted of processing raw ore and scrap tungsten concentrates to produce ammonium paratungstate (APT) and subsequently formulating APT to metal tungsten powder and tungsten carbide powder. Other specialty products including tungsten carbide powder plus cobalt and other material for plasma spraying; tungsten titanium carbide powder; tantalum carbide powder; tungsten spray powder; crystalline tungsten powder; and molybdenum spray powder were also produced.
1948	National Reconditioning Corporation changes its name to Wah Chang Smelting and Refining Corporation (WCSRC).
1948 - 1964	Site operated by WCSRC.
1964	WCSRC leases equipment/property to the Wah Chang Corporation (WCC) which continued to operate the facility.
April 1967 - 1972	Teledyne acquired the stock of WCC and the two companies merged. Operations at the site continued by Teledyne-Wah Chang Corporation.
1972	WCSRC formed a wholly owned subsidiary (Li Tungsten Corporation) which operated the facility until filing for bankruptcy in 1985.
1984	Property acquired by the Glen Cove Development Company (GCDC). GCDC is a general partnership duly organized and existing under the laws of the State of New York and is owned by the Old Court Holdings Company and the Old Court Joint Ventures, Inc., both of which, in turn, are wholly-owned subsidiaries of Old Court Savings and Loan, Inc., (in Receivership) located in Maryland

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TABLE 2-1 (continued) CHRONOLOGY OF EVENTS AND SITE INVESTIGATIONS I i Tungsten Site Glen Cove, New York

DATE	EVENT
1984	GCDC continues to lease the site to Li Tungsten Corporation. Market for tungsten in decline.
June 1985	Li Tungsten Corporation files for bankruptcy. Manufacturing operations at the facility cease.
May 1988	RTP Environmental Associates, Inc., (Westbury, NY) completes Site Investigation Report for Campon Reality Corporation (RTP, 1988). Site investigation undertaken to evaluate environmental conditions prior to residential development. Geraghty and Miller was subcontracted to perform the hydrogeology investigation.
March 29, 1989	New York State Department of Environmental Conservation (NYSDEC) performs site inspection.
April 14-16, 1989	USEPA assumes lead enforcement role on response actions at the site. USEPA FIT2 contractor (NUS) initiates Preliminary Assessment.
July 21, 1989	Administrative Order On Consent (AOC) issued by USEPA to Glen Cove Development Corporation which specified nine (9) interim remedial actions.
September 18, 1989	USEPA FIT2 contractor (NUS) issues Preliminary Assessment Report (NUS, 1989).
April 4, 1990	Interim remedial actions completed and final report submitted (HART, 1990).
September 28, 1990 :	USEPA FIT2 contractor (NUS) issues Site Inspection Report (NUS, 1990).
July 1991	Li Tungsten site proposed for inclusion on the National Priorities List (NPL).
February 12, 1992	Special Notice letters were sent by USEPA to five PRPs (Teledyne, Inc.; Li Tungsten Inc.; the Glen Cove Development Corporation; Wah Chang Smelting and Refining Corporation; and Mr. John Li (son

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TABLE 2-1 (continued) CHRONOLOGY OF EVENTS AND SITE INVESTIGATIONS Li Tungsten Site Glen Cove, New York

DATE EVENT

of Mr. K. C. Li). These letters solicited the involvement of the PRPs in the investigation of the site.

August 26, 1992

Malcolm Pirnie receives work assignment to prepare

RI/FS Work Plan.

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separation; roasting; leaching; floatation; and fusion. A generalized flow sheet of the treatment processes is show in Figure 2-4.

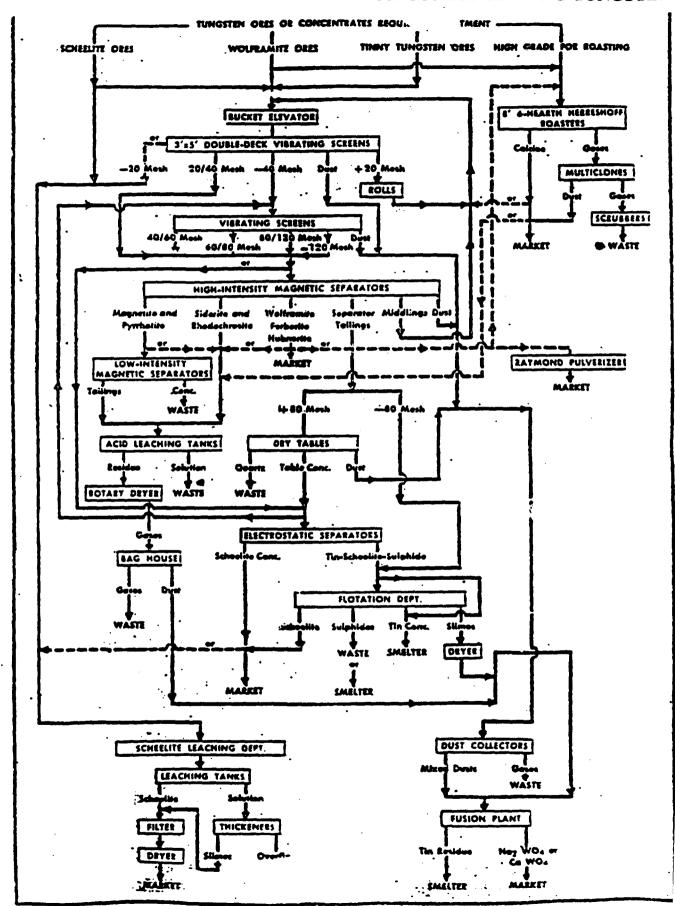
Numerous aboveground wooden, steel or fiberglass tanks were used in performing some of these treatments and to store reactants (e.g., hydrochloric acid, ammonia, hydrogen) and/or intermediate compounds (e.g., APT). Many of these tanks still contain some hazardous and inorganic liquids. As the tungsten ore moved through its various processing stages, the naturally occurring radioactive isotopes of thorium, uranium, and radium became more concentrated in the residual waste or slag. There are indications that some of this slag was placed in waste piles at the ground surface and buried on site (NUS, 1989; 1990). Accessory metals which constitute the impurities that were removed during the extraction process include: antimony, arsenic, barium, bismuth, copper, cobalt, chromium, lead, manganese, mercury, molybdenum, nickel, thorium, uranium, vanadium, and zinc.

In 1948 the National Reconditioning Company changed its name to Wah Chang Smelting and Refining Corporation (WCSRC). WCSRC continued to operate the site until 1964 when they leased the equipment and property to Wah Chang Corporation (WCC). In 1966 Teledyne acquired the stock of WCC and the two companies merged. Operations at the site continued by Teledyne-Wah Chang Corporation.

In 1972 WCSRC, which had been leasing the equipment and property to Teledyne-Wah Chang Corporation, formed a wholly owned subsidiary (Li Tungsten Corporation) which continued to operate the facility. In 1984 the property was acquired by GCDC. GCDC is a general partnership duly organized and existing under the laws of the State of New York and is owned by the Old Court Holdings Company and the Old Court Joint Ventures, Inc., both of which, in turn, are wholly-owned subsidiaries of Old Court Savings and Loan, Inc., (in Receivership) located in Maryland. GCDC continued to lease the site to Li Tungsten Corporation until 1985 when Li Tungsten Corporation ceased operations at the site and filed for bankruptcy.

There is very little specific documented knowledge on waste volumes that were generated or waste disposal practices. Drummed waste is also reported to have been buried on-site in a portion of Parcel B (NUS, 1989, 1990). Liquid wastes are believed to have been disposed of through numerous subsurface drainage pipes that have been noted in the

FLOWSHEET OF WAH CHANG'S PLANT FOR CONCENTRATING TUNGSTEN



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bulkhead and empty directly in Glen Cove Creak. State Pollution Discharge Elimination System (SPDES) permits allowed for up to as many as 250,000 gallons per day of discharge to Glen Cove Creek. Mud Pond and the two Mud Holes were also reportedly used to dispose of liquid wastes.

On April 14, 1989 the USEPA received a request from the NYSDEC to use its Superfund authority to respond to threats posed by hazardous materials at the site. USEPA's preliminary assessment and site inspection of site conditions (NUS, 1989; 1990), revealed a large quantity of slag which was emitting low-level beta-gamma radiation. In addition, large quantities of laboratory reagents, various hazardous materials in drums and tanks, asbestos, transformers, and cylinders containing compressed liquids and gases were found in several buildings. Air monitoring showed no dangerous levels of organic compounds either on site or off-site. As a result of the conditions identified at the site, the USEPA issued an Administrative Order on Consent (AOC) to GCDC to stabilize all potential threats to the public and the environment.

Fred C. Hart Associates, Inc., (HART) was hired by GCDC to coordinate the nine (9) interim remedial actions identified in the AOC (HART, 1990). Additional removal/remedial actions were also undertaken by GCDC. A list of the interim remedial actions and the additional actions completed at the site is summarized in Table 2-2.

The Hazard Ranking Score (HRS) for the Li Tungsten site was 50.00 which is above the 28.5 threshold value for inclusion on the NPL (NUS, 1991). In July 1991 the Li Tungsten site was proposed for inclusion on the NPL and in October 1992, the site was placed on the NPL.

2.3 Current Conditions

The Li Tungsten site ceased operations in June 1985 and has been inactive since. Site security (fencing and guard) was addressed as one of the interim remedial measures in the AOC. Although a security guard is present on-site 24 hours a day, the site could be entered without knowledge of the security guard through breaks in the fence. During the site visits, observations were made that trespassing has occurred. Many of the salvageable fixtures

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TABLE 2-2 SUMMARY OF INTERIM REMEDIAL ACTIONS AND ADDITIONAL REMOVAL ACTIONS Li Tungsten Site Glen Cove, New York

AOC Specified Tasks

The AOC contained a schedule for completion of the nine tasks listed below. There is insufficient information available to determine exactly when these activities were completed, but generally they occurred between the date the AOC was signed on July 21, 1989 and the date the Remedial Action Report was issued on April 4, 1990.

Site Security

Repairs were made to all existing fences and gates. New fence was installed in two areas. All gates were made functional and fitted with locks.

Radioactive Materials

- Twelve (12) drums (or 113 cubic feet) of equipment, thorium metal and other materials (HART, 1990, p.13), plus a small furnace were removed by NDL on December 11, 1989.
- Twenty (20) yards of radioactive process ore slag was relocated to a secure area within the Dice building (HART, 1990, p.13).

Laboratory Chemicals

- Fifty-two (52) 55-gallon and 80-gallon overpacks and twenty (20) 5-gallon pails of labeled laboratory chemicals were prepared for shipment to Cycle Chem.
- Eight (8) 55-gallon drums of unknown liquid laboratory chemicals were placed in the staging area.
- One (1) 55-gallon drum of unknown solid laboratory chemicals were placed in the staging area.

Drummed Chemicals

• The liquids from approximately 150 - 200 unknown drums were bulked for removal and disposal (HART, 1990, p24).

Tank Characterization

- A total of 223 tanks were identified on the three site parcels [A 197 tanks (112 empty); B 6 tanks (all empty); and C 20 tanks(14 empty)] (HART, 1990, p. 35).
- Disposal of tanks was not specified as part of IRA (HART, 1990, p.24).

TABLE 2-2 (continued) SUMMARY OF INTERIM REMEDIAL ACTIONS AND ADDITIONAL REMOVAL ACTIONS Li Tungsten Site Glen Cove, New York

Asbestos Sampling

- Five (5) high volume air samples were collected (Carbide Building; West Dice Building; Loung Building; Dickson Warehouse; and Benbow Building) and analyzed by transmission electron microscopy (TEM) with no indication of airborne asbestos (HART, 1990, p. 57).
- Fifty-one bulk samples were collected from Parcels A and C and analyzed by polarized light microscopy (PLM). Slightly more than half of the samples (53%) reflected the presence of asbestos containing materials (ACM). Results are presented in Plate 2 (HART, 1990).

Creek Sediments

Five (5) sediment samples were collected from Glen Cove Creek and two (2) sediment samples and two (2) sediment core samples were collected from Hempstead Harbor. No enhanced levels of radionuclides were detected in the creek or the harbor.

Transformer Characterization

- Thirty eight (38) samples were collected from transformers or other electrical equipment. Eleven (11) samples collected reflected concentrations of PCBs greater than 50 ppm; three (3) units reflected concentrations greater than 500 ppm.
- Although not specified in the AOC, transformer oils were drained from all units; some were disposed of as PCB oils, others as non-PCB oils. The carcasses of three (3) transformers were also disposed of as PCB solids (HART, 1990, p.68).

Mercury Spill Cleanup

• An area inside the Benbow Building (Parcel C) was identified as having mercury on the floor. A commercially available mercury absorbing salt was used to absorb the mercury. Portions of the subfloor conduit which contained mercury could not be cleaned up because heavy equipment that was present made the area inaccessible. The room was boarded up and labelled to indicate the presence of residual mercury.

Additional Tasks Not Specified in AOC

The additional tasks listed below were completed by GCDC and with the concurrence of USEPA either prior to issuance of the AOC or concurrent with the AOC specified tasks listed above. A separate order was issued in April 1989 for the removal of the anhydrous ammonia. In general, these tasks were completed between June 1989 and April 1990.

TABLE 2-2 (continued) SUMMARY OF INTERIM REMEDIAL ACTIONS AND ADDITIONAL REMOVAL ACTIONS Li Tungsten Site Glen Cove, New York

Pressurized Cylinders

- Twenty-six (26) cylinders were identified for removal. Twenty-four (24) of these cylinders were clearly marked with the name of the owner/distributor. The owners/distributors were contacted and the cylinders were removed.
- Two (2) cylinders remain at the site their contents are unknown. They were scheduled for sampling and analysis in April 1990. The results of this sampling is not known.

Additional Laboratory Overpacks

Due to the number of chemicals (over 2500 individual containers; 500 with labels) found in the laboratories, offices, storage spaces in Parcel A, strict adherence to the limitation of the interim remedial action (200 laboratory chemicals) would have left a large quantity of chemicals on-site. Additional chemicals were removed, however, some may still remain

Radioactive Slag Relocation

• Three (3) dump truck loads (approximately 20 cubic yards) of radioactive slag were moved from Parcel A (near the fence at Herb Hill Road and Garvies Point Road) to inside the West Dice Building. The slag was placed on pallets, covered with plastic, and labeled with signs indicating a radioactive hazard.

Anhydrous Ammonia Removal

 One (1) tank of anhydrous ammonia on Parcel A was emptied pursuant to a separate order issued in April 1989. The anhydrous ammonia was removed and returned to its distributor (HART, 1990, p. 69).

Methyl Ethyl Ketone Peroxide (MEKP) Removal

• One (1) pint of MEKP was removed from the refrigerator in the main office building (dark room) for disposal (HART, 1990, p. 70).

Air Sampling

- Inorganic Acid Gases fluoride was found in excess of one field blank and was thought to be due to hydrofluoric acid found in several drums.
- Volatile Organic Compounds not detected in significant quantities.
- Inorganics all samples were significantly below ACGIH published Threshold Limit Values (TLVs). No difference was found between air samples collected inside the buildings and those collected outside.

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(e.g., copper wiring and piping) have been removed and general debris (e.g., washing machines, mattresses) have been left behind.

to contain elevated radionuclide concentrations.

3.1.7 Characteristics of Chemical Contamination

Characteristics of chemical contamination on the site stem from activities associated with the production of tungsten carbide powder, tungsten wire and welding rods. To produce these products, monazite sand and tungsten ore or concentrates were smelted between the 1942 and 1985. The treatment processes used to extract tungsten metal from these materials generated a residual slag (waste ore) which tended to concentrate radioactive isotopes of uranium, thorium and radium, and other heavy metal impurities. The slag, as well as some processed and unprocessed, ore was stored on-site in wooden crates, piles, and drums. Much of this material still remains on the site and some of it is believed to have been disposed of on site (Parcels B and C).

Potential contaminants on the site include commercially prepared strong acids, strong bases, organic solvents, aqueous ammonia, mercury and cyanide which were used in the treatment processes. The acids were used for leaching of impurities out of the tungsten where mechanical separation was not effective. An on-site laboratory also existed where the tungsten product was analyzed for impurities and either sent for reprocessing or identified as a finished product. The majority of chemicals used in the laboratory were removed as part of the interim remedial actions (HART, 1990). Other organics used on the site included PCBs in transformers, and fuel oil which was stored in several tanks, including one 500,000 gallon aboveground storage tank.

Asbestos containing materials (ACM) has been found on-site in siding shingles, roof tiles, tank covers and pipe insulation. ACM has also been found on the ground at the site.

3.1.8 Sources and Distribution of Contamination

As described earlier, several investigations have been completed at the site (RTP, 1988; G&M, 1988; NDL, 1989; HART, 1990; NUS, 1989; 1990). The results of these investigations were used to prepare the following sections which summarize the current understanding of environmental conditions at the site.

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The primary sources of contamination on the site include processed and partially processed tungsten ore present in drums, wooden crates, and piles located both inside and outside the buildings. Removal of these source materials is proposed in this Work Plan as an interim remedial action prior to initiating the RI field investigation. Other potential sources include; Mud Pond and two Mud Holes which were used for disposal of wastewater; the disposal area located on Parcel B; unconfirmed disposal areas on Parcel C that is devoid of vegetation; underground storage tanks (whose locations and contents are unknown), and a 500,000 gallon aboveground fuel oil tank.

Secondary sources of contamination include the on-site soil; off-site groundwater from the Mattiace property, the former Powers-Chemco property and/or a former dry cleaner; and the storm and process drains on-site and off-site. Removal of asbestos is also proposed in this Work Plan as an interim remedial action prior to initiating RI field investigation (See Section 3.7).

Chemical contamination is distributed throughout the groundwater, surface water, soils and sediments at the site. Volatile organic compounds in the groundwater may originate from off-site sources, including a former dry cleaning establishment to the east of Parcel B and the Mattiace site (NUS, 1990). No on-site source of organic contaminants has been identified. The predominant contamination attributable to on-site sources is inorganic metals. Inorganic metals are found at the majority of the groundwater sampling locations. Inorganic metals have been identified in the on-site surface water and sediment contamination, including Mud Pond, the Mud Holes, the pond and associated drainage stream on Parcel B, the standing water in the building, and open tanks.

3.1.8.1 Chemical Characteristics of Soil

This section presents a summary of the chemical characterizations of the soils based on existing data (NUS, 1990). Soil samples were collected at a total of 10 locations (S-1 through S-10) as shown on Figure 3-1. The samples were analyzed for volatile and semi-volatile organic compounds, pesticides/PCBs, and inorganic compounds (metals and non-metals).

REFERENCE 8

Pursuant to the Labor Law and Industrial Code Rule No. 38, and in reliance on statements and representations heretofore made by the licensee designated below, a license is hereby issued authorizing such licensee to transfer, receive, possess and use the radioactive material(s) designated below; and to use such radioactive materials for the purpose(s) and at the place(s) designated below. This license is subject to all applicable rules, regulations, and orders now or hereafter in effect of all appropriate regulatory agencies and to any conditions specified below.

Licensee		3. License nur	nber
1. Name Wash Chang Smelting and Refining Company of inerica, Inc. 2. Address 63 Herb Hill Road Glen Cove, New York		4. Expiration date Valid until terminated	
6. Radioactive materials (element and mass number) 1. Thorium 2. Thorium	7. Chemical and/or physical form 1. Thorium oxide 2. Thorium nitrate		8. Maximum quantity licensee may possess at any one time 1. 2300 pounds 2. 750 pounds
			Total thorium not to exceed 15.4 curies 15 6 (sum un calculation) AN

CONDITIONS

- 9. Authorized use. (Unless otherwise specified, the authorized place of use is the licensee's address stated in Item 2 above.)1. As insulator in vacuum furnace.
 - 2. Production of theriated tungston newder as step in manufacturing of welding rods.
- 10. The licensee shall conduct operations involving the use of sources of radiation in compliance with the requirements of New York State Industrial Gode Rule Fo. 38, "Padiation Protection".
- 11. Iny disposal of radioactive waste by the licenses by burial, through the sanitary sever, or by other release to the environment shall be in accordance with the provisions of Part 16, New York State Sanitary Code Records of all such disposal shall be naintained by the licensee. Monitoring procedures shall be instituted where necessary to demonstrate that concentrations and quantities of radioactive raterial so disposed of do not exceed permissible levels.

 12. The agreement material described in Items 6, 7 and 3 above:
 - A. Shall be used only by or under the supervision of either A. Horra or A. Bathie
 - B. Shall not be used in or on human beings, in products intended for uncontrolled distribution to the general public, nor in field applications where radioactivity is released.

	FOR THE NEW YORK STATE DEPARTMENT OF LABOR
Date	by

Form CCL-62SL (8-63)

STATE OF NEW YORK RADIOACTIVE MATERIALS LICENSE

Page 2	_ct	Pages
_		NF B
		2 of 2

License Number 743-0464

Reference Number: 1

- C. Shall be possessed and used by the licensee in accordance with statements, representations and procedures contained in his application dated February 26, 1964, and in related documents as follows:
 - 1. Letter to the United States Atomic Energy Commission dated February 20, 1961, signed by Allen Lau.
 - 2. So much of Part 40, Title 10, Gode of Federal Regulations as is applicable to operations of the licensee and not in conflict with Code Rule 33 or the other conditions of this license.

10360

Date March 19, 1964

APA:10

Form COL-66SL (8-63)

FOR THE NEW YORK STATE DEPARTMENT OF LABOR

Mathan Solemon, Ph.D., M.D.

Chief, Radiological Health Unit For: Morris Kleinfeld, Director, DIE **REFERENCE 9**



UNITED STATES

ATOMIC ENERGY COMMISSION

WASHINGTON 23, D. C.

LO-SL3 CALIPOS

> Web Chang Swelting & Refining Company of America, Inc.

Woolmerth Building

Attention: Er, Allen Las, Ass't Treasurer

SOURCE MATERIAL LICENSE

License No. D-607

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Dated: DEC 6 1957

Contlemen

Pursuant to the Atomic Energy Act of 1994 and Section 40.21 of the Code of Federal Regulations, Title 10 - Atomic Energy, Chapter 11 Part 40 - Control of Source Material, you are hereby licensed to receive possession of and title to sixteen hundred (1600) possess of wrendow and thorium country that term of this license, for monter an analytical

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You are further licensed to transfer and deliver possession of and title to refined source material to any person licensed by the Atomic Energy Commission, within the limits of his license and light

As a condition of this license; you are required to maintain records of your inventories; receipts and transfers of reclined source material.

This licenses is subject to all the provisions of the Aconte Energy Act of 1954 now on hereafter, in effect and to all value rules and regulations of the U.S. Atomic Energy Commission, including 10 CFR 20, Standards For Protection Against Rediction.

Beither, this license nor any right under this license shall be assigned or otherwise transferred in violation of the provisions of the Atomic Energy Act of 1954.

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As a condition of this license, you are required to maintain records of your inventories; receipts and transfers of refined source material.

This license is subject to all the provisions of the Atomic Energy Act of 1954 now or hereafter in effect and to all the lideriles and regulations of the U.S. Atomic Energy Commission, including 10 (1872) "Standards For Protection Against Radiation."

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REFERENCE 10

RADIOLOGICAL SURVEY PHASE II INVESTIGATION GARVIES POINT GLEN COVE, NEW YORK

DUPLICATE 6/5/10

Prepared by:

Fred C. Hart Associates, Inc. 470 Park Avenue South 11th Floor New York, New York 10016

and

The NDL Organization, Inc. P.O. Box 791 Peekskill, New York 10566

June 5, 1990

TABLE OF CONTENTS

<u>Section</u>	<u> Title</u>	Page
1.0	INTRODUCTION	1
2.0	RADIOLOGICAL SURVEYS 2.1 Summary of Phase I Investigation 2.2 Phase II Aerial Photograph Review 2.2.1 Objective 2.2.2 Site History	2 2 6 6 6
	2.2.3 Fill Distribution and Thickness	7 7 7 7
	2.4 Phase II Subsurface Investigation	21 21 21 25
3.0	DISCUSSION	28 28 28 31

LIST OF APPENDICES

Appendix A - Calculation of Effective Depth of Ore Detection

Appendix B - Trench Logs

Appendix C - Laboratory Analytical Data

LIST OF FIGURES

Number	<u>Title</u>	Page
2-1	Approximate Sample Locations in Phase I	3
2-2	Fill Thickness Map	8
2-3	Approximate Areas of Elevated Gamma Fluxes from the 0-6 Foot Interval	20
2-4	Trench Locations	22
3-1	Estimated Extent of Elevated Levels of Radionuclide Concentrations	30
3–2	Revised Health and Safety Plan for Remedial Investigation	32

LIST OF TABLES

<u>Number</u>	<u> Title</u>	Page
2-1	Garvies Point Phase I Radiological Survey Analytical Results	5
2-2	Large Area Gamma Ray Survey Results, January 23 - February 8, 1990	11
2-3	List of Grid Points That Exceed Background Levels	19
2-4	Correlation between Gamma Ray Flux Measurements and Trench Locations	23
2-5	Summary of Radionuclide Concentrations in Trench Samples	26

1.0 INTRODUCTION

Environmental investigations at the Li Tungsten facility in Glen Cove, New York, have indicated that residual ore at the facility contains naturally occurring radionuclides such as thorium and uranium. These radionuclides are present in the ore naturally and were concentrated in the residual ore by the refining process. In April 1989, Fred C. Hart Associates, Inc. (HART) was notified of unsubstantiated allegations that some of the residual ore from the Li Tungsten operations may have been deposited at the nearby Garvies Point Condominium site, also in Glen Cove, while this site was operated as a municipal landfill. At the time of these allegations, HART was in the process of finalizing a Remedial Investigation Work Plan for the Garvies Point site for approval by the New York State Department of Environmental Conservation (NYSDEC).

In order to investigate these allegations, HART conducted a preliminary radiological survey (referred to as the Phase I Investigation in this document) at Garvies Point. The scope of this survey was described in the Radiological Survey Work Plan submitted to NYSDEC on June 1, 1989. The survey was conducted on a 100 by 100 foot system of grid points with a microR meter. This instrument measures real time radioactivity from the surface to a maximum depth of two feet. Samples were subsequently collected from three locations at the site where elevated readings were noted in the microR meter survey. HART prepared a report summarizing the results of the Phase I survey which was submitted to NYSDEC on November 27, 1989. For completeness, the results of the survey are also summarized in Section 2.1 in this report.

Based upon the results of the Phase I survey, a Phase II Radiological Survey was proposed by HART. The Phase II survey was designed to provide more detailed information on the horizontal and vertical extent of radionuclides at Garvies Point. The scope of the Phase II survey was outlined in the Radiological Survey Results report of November 23, 1989 and consisted of an aerial photograph review; a large area gamma ray survey on a 50 by 50 foot grid with an instrument capable of penetrating up to six feet of soil; and the excavation of trenches in background areas

and areas of elevated gamma ray readings. The Phase II investigation was implemented at the site between January 23 and February 13, 1990. The results of the survey are provided in Sections 2.2 - 2.4 of this report.

2.0 RADIOLOGICAL SURVEYS

2.1 Summary of Phase I Investigation

A field team consisting of two HART personnel and a certified health physicist performed the radiological survey on August 23, 1989. Initially, a 100 by 100 foot grid was established at the site by Baldwin and Cornelius, P.C. and HART personnel. Data was collected along the grid with two microR meters from two different manufacturers in order to verify measurements. A Geiger counter was not used because this instrument was not thought to be sensitive enough to measure the levels of radiation that were anticipated. The level of radiation was measured at each gridpoint at the ground surface and one meter above the surface. Any elevated readings between the grid points were also noted. The measurements and all relevant observations were recorded in a bound field notebook.

Agreement between the two different microR meters was excellent, indicating that the on-site measurements were accurate. Most of the site had radiation levels between 3 and 15 uR/hr as measured by the microR meters. These levels are within the normal background range of up to 20 uR/hr.

Three areas with measurements that exceeded site background were noted; the locations of these areas are shown in Figure 2-1. The area near the driveway (Area 1) had readings between 20 and 25 uR/hr. Although most of this area was covered with high grass, the highest measurements occurred in a 4 foot by 10 foot area of unvegetated soil. When digging below the surface in this area, readings up to 50 uR/hr were recorded at a depth of 6 to 18 inches. A sample for laboratory analysis was collected from this interval.

Areas 2 and 3 were near each other but were not contiguous. Readings up to 60 uR/hr were recorded in Area 2 while Area 3 had readings up to 30 uR/hr. The elevated readings occurred in an approximately 10 foot diameter region at Area 2 and in an approximately 6 foot diameter region at Area 3. Similar to Area 1, readings increased below the surface in these two areas. A fairly discrete reddish clay-rich sand layer which had readings of 125 to 140 uR/hr was found in Area 2 at 6 to 10 inches below the surface. A sample for laboratory analysis was collected from this discrete clay layer. Readings at Area 3 increased from 30 uR/hr at the surface to 40 to 60 uR/hr about 10 inches below the surface. The entire interval was sampled for laboratory analysis.

In addition to the three samples collected from areas where above background levels of radionuclides were detected, one sample was collected at a grid point with background radiation levels for comparison. All samples were obtained by digging below the surface with a spade and filling a one liter glass jar. All samples were packaged in a cardboard box with styrofoam packing material and shipped to Recra Environmental, Inc. for analysis. The outside of the package was scanned with the microR meter at the time of shipment and no measurements above background were obtained. All soil samples were analyzed for gross alpha and gross beta radioactivity and the gamma spectrum of each sample was determined using a germanium detector.

The results of the radiological analyses are summarized in Table 2-1. The levels of radioactivity measured in the samples correlated qualitatively with the field measurements; i.e., the background sample had the lowest levels, and Area 2 had the highest levels.

Based upon the results of the Phase I radiological survey and the soil sample analyses, a Phase II radiological survey was proposed for the site. The purpose of the Phase II survey was to further characterize the vertical and lateral extent of radioactive materials. A smaller grid configuration and a different type of survey instrument were used to provide more detailed data. The Phase II survey included three tasks: an aerial photograph review, a large area gamma ray survey and a subsurface investigation.

(2529n-4)

TABLE 2-1

GARVIES POINT PHASE I RADIOLOGICAL SURVEY
ANALYTICAL RESULTS

1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1								Gamma Spectra (pCi/g)						
Samole	Depth	(uR/hr)	(PCI/a)	(PCi/a)	11-208	Pb-212	81-212	Pb-214	Bi-214	<u>K-40</u>	Ac-228	<u>Ih-227</u>	Ih-234	<u>U-235</u>
Background	Surface	3-15	4:8 <u>+</u> 2.8	12±4	0.5 <u>+</u> 0.1	1.1 <u>+</u> 0.2	1.3 <u>±</u> 0.2	1.2 <u>+</u> 0.2	1.1 <u>±</u> 0.2	15 <u>+</u> 2	2.8±0.3	0.8±0.3	2.8±0.8	0.3 <u>±</u> 0.
Area 1	6-8 in	50	25 <u>+</u> 6	28 <u>±</u> 5	1.9±0.2	4.9 <u>+</u> 0.5	8.7 <u>±</u> 0.9	10 <u>±</u> 1	8.0±0.8	15 <u>±</u> 2	13 <u>+</u> 2	2.6 <u>±</u> 0.7	19 <u>±</u> 2	1.2 <u>4</u> 0.7
Area 2	6-10 in	125-140	580 <u>±</u> 60	520 <u>1</u> 60	80 <u>+</u> 8	210±30	140±20	51 <u>+</u> 6	4416	28±3	490±50	10±1	250±30	7.0 <u>±</u> 0.
Area 3	0-10 in	30-60	200 <u>±</u> 20	140 <u>+</u> 20	8.4 <u>+</u> 0.9	23 <u>±</u> 3	15 <u>±</u> 2	51 <u>±</u> 6	41 <u>±</u> 5	16±3	51 <u>±</u> 6	12 <u>+</u> 2	100 <u>±</u> 10	7.1 <u>±</u> 0.8

2.2 Phase II Aerial Photograph Review

- 2.2.1 Objective. The photographs were reviewed in order to determine the extent of landfilling at the site. This information was used to decide which areas of the site were to be investigated in greater detail during the large area gamma ray detector survey and trenching tasks.
- 2.2.2 <u>Site History</u>. The aerial photographs can be grouped into three periods, each of which is characterized by a different use of the Garvies Point Site. The earliest period is shown in the photographs taken in 1950 and 1955. During this time, the site was relatively dormant. Small boats and a building are visible near the southwest part of the site. There are several small trenches which appear to have been installed to facilitate drainage to Glen Cove Creek. Overall, the topography of the site does not appear to have been significantly altered by landfilling or other site activities. Although the U.S. Army Corps of Engineers (USACE) dredged Glen Cove Creek in 1933, 1934, and 1948, available records do not state whether this material was disposed at the site or elsewhere.

The first aerial photograph in which landfilling activities at Garvies Point are apparent is the 1962 photograph. However, since there are no aerial photographs from the period between 1955 and 1962, it is possible that landfilling activities started prior to 1962. Landfilling activities are also apparent in the photographs taken in 1966, 1969 and 1972. The most obvious change is the filling of a tidal embayment in the eastern part of the site. Other filling took place in the center of the site immediately north of the tidal flat, and east of the beach at the western end of the site. Dredge spoils from Glen Cove Creek are known to have been disposed of at the site by USACE in 1960 and 1965. The bulkhead at the mouth of Glen Cove Creek at the western end of the site was built between 1966 and 1969, although it may not have been its present height at that time. A photograph taken in 1978 appears to represent a period during which landfilling activities had stopped and vegetation was allowed to grow.

The most recent period of use is characterized by preparation for, and the start of, development of the site for use as a residential area. This (2529n-6)

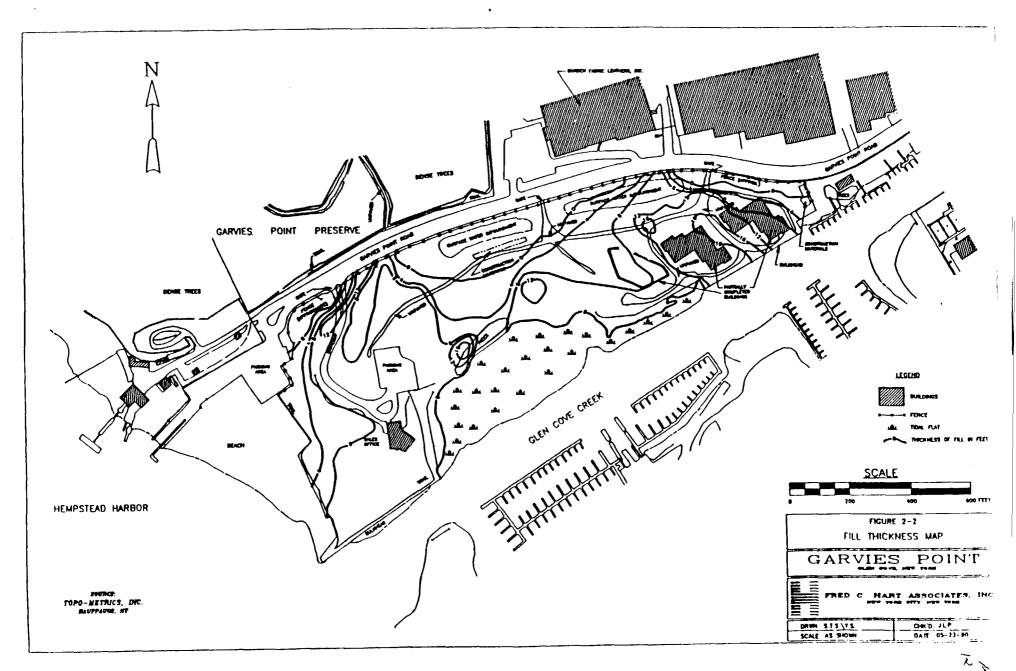
use is illustrated in the photographs taken in 1984, 1986 and 1989. During this time, the major site features which are visible today were built. These include the sales office and driveway, surface water retention ponds, and the bulkhead and partially completed buildings at the eastern end of the site.

2.2.3 <u>Fill Distribution and Thickness</u>. The extent and thickness of the fill at Garvies Point can be estimated by comparing the present topography of the site with the topography of the site prior to landfilling. For purposes of this study, the aerial photograph taken in 1950 was assumed to represent the original site topography. The present topography is shown in a topographic map made from the aerial photograph taken in 1989. An estimate of the fill thickness based on these data is shown in Figure 2-2.

Because there are few points on the 1950 aerial photograph with known elevations, the fill thicknesses shown in Figure 2-2 are approximate. Furthermore, there is no distinction in the figure between different fill materials or different periods of landfilling activities. Given these qualifiers, it is still clear that much of the site has been landfilled. Fill materials are thickest in the former area of the tidal embayment on the eastern end of the site, where they reach approximately 16 feet in thickness. The tidal flat and some areas along Garvies Point Road have not been filled.

2.3 Phase II Large Area Gamma Ray Survey

- 2.3.1 Objective. As the depth penetration of the microR meter used in the first survey was only one to two feet and fill thicknesses were thought to be 10 to 15 feet, the large area gamma ray survey was conducted to determine whether or not above background levels of radiation existed at depths of up to six feet below the surface. The results of this survey were also used in the selection of trench locations in areas of elevated gamma ray fluxes and in background locations.
- 2.3.2 <u>Methodologies</u>. The large area gamma ray survey took place from January 23 to February 8, 1990. The survey was performed by the NDL (2529n-7)



Organization, Inc. (NDL) of Peekskill, NY and was overseen by HART personnel. The large area gamma ray detector used in the survey was built by Ted Rahon of NDL.

The gamma ray detector consisted of an eight inch diameter, two inch thick. Cesium Iodide (CsI) scintillation crystal. It has been termed "large area" because of its 50 square inch face (324 sq.cm.) as opposed to the usual 0.2 to 0.8 square inch face of Sodium Iodide (NaI) detectors used in microR meters. The two inch crystal thickness and the higher gamma ray absorption coefficient of CsI make the detector more sensitive to high energy gamma rays than a microR meter. A single channel analyzer was used with the detector so that only gamma rays in the 2.6 MeV energy region were counted. The analyzer threshold was set so that the system did not respond to Cs-137 (0.662 MeV) or Ra-226 (0.609 MeV. 1.76 MeV) fields. This threshold setting made the detector effectively unresponsive to all naturally occurring radionuclides except T1-208 (2.6 MeV). Even if "clean" overlving soil had elevated U-238 series concentrations, it would not effect the sensitivity of the system to detect the T1-208 gamma ray.

The upper level discriminator of the analyzer was used to reduce detector background from cosmic radiation. The upper level was set such that the count rate from a natural thorium source was not significantly affected by opening or closing the window. However, the high energy background (>3 MeV), due mainly to cosmic rays, was substantially reduced. To limit counting error, sufficient counts at each location were collected to yield less than a 5% error. Thus, the counting period used at each grid location was determined by the count rates encountered. Counting periods ranged from 2 minutes at grid points with elevated gamma ray counts to 10 minutes at the background grid points.

Measurements on undisturbed, native Garvies Point soil indicated that indigenous thorium concentrations were very low and would provide the desired low background for the survey. With these background conditions and the instrument setup described above, a truck-load size mass of tungsten ore with a natural thorium concentration of 100 pCi/g could be

detected under 1 to 2 meters of soil with a density of 1 to 1.5 g/cc (see calculations in Appendix A). The large area gamma ray detector was tested at New York University Medical Center AJ Lanza Laboratories using a National Bureau of Standards natural thorium source and sand as an attenuator prior to performing the survey at Garvies Point.

The gamma ray survey at the site was conducted on a 50 foot by 50 foot grid, offset by 25 feet from the grid stakes. This grid configuration was used to avoid retesting areas surveyed with the microR meter in the Phase I investigation. At each survey location, the grid point, gamma ray count and length of time the detector was run were recorded in a bound field notebook. As a result of buildings or surface water, several grid locations were offset to make them accessible.

2.3.3 Results. The data collected from the large area gamma ray survey is summarized in Table 2-2. In the field, the number of counts per minute (cpm) was recorded for each location. The criteria used to determine if a reading should be classified as "elevated" was two standard deviations above the local background. This criteria was selected because it ensured that 95% of the data selected would be above background and it minimized the chances of missing areas that were truly above background. Only a few of the gamma ray readings fell between two and three standard deviations of background, so the move conservative criteria of two standard deviations was applied. After 40 to 50 points in one section of the property were measured, the mean and standard deviation were calculated for readings collected from locations thought to be free of radioactive material. For example, the first 40 points measured on the western side of the property, excluding Area 1, resulted in a mean count rate of 125 cpm and a standard deviation of 24 cpm. Thus, the background level for the west side was 173 cpm (125+2x24). The background levels in both the middle and eastern sections of the property were approximately 145 cpm. A total of 25 grid points exceeded the background levels defined as local background plus two standard deviations. A list of these grid points is shown in Table 2-3 and the locations of these points are shown in Figure 2-3.

TABLE 2-2

LARGE AREA GAMMA RAY SURVEY RESULTS

JANUARY 23 - FEBRUARY 8. 1990

Location	Counts Per Minute	uR/hr	Location	Counts Per Minute	uR/hr
Z-10, 1	111± 5	 5	Z+25, 1	125± 4	6
Z+75, 1+25	127± 4	6	Z+75, 1+75	104± 3	6
Z+75, 2+25	194± 4*	8	Z+75, 2+75	112± 3	7
Z+75, 3+25	98± 3	7	Z+75, 3+75	120± 3	7
Z+75, 4+25	208± 6*	9	Z+75, 4+75	106± 3	7
Z+75, 5+25	129± 4	8	Z+75, 5+75	106± 3	8
Z+75, 6+25	103± 3	7	Z+75, 6+75	118± 3	8
A+25, 1+25	114 <u>+</u> 3	6	A+25, 1+75	106± 3	7
A+25, 2+25	126± 4	7	A+25, 2+75	132 <u>+</u> 4	7
A+25, 3+25	126± 3	7	A+25, 3+75	120± 3	8
A+25, 4+0	140± 4	8	A+25, 4+75	118 <u>+</u> 3	8
A+25, 5+25	122± 3	9	A+25, 5+75	148± 4	10
A+25, 6+25	139± 4	11	A+25, 6+75	131± 4	9
A+25, 7+25	133 <u>+</u> 4	9	A+25, 7+75	196± 6*	14
A+75, 1+25	126± 4	7	A+75, 1+75	135± 4	7
A+75, 2+25	114± 3	7	A+75, 2+75	129 <u>+</u> 4	7
A+75, 3+25	130± 4	: 8	A+75, 4+0	164± 4	11
A+75, 4+25	145± 4	9	A+75, 4+75	126± 4	9
A+75, 5+25	115± 3	8	A+75, 5+75	96± 3	6
A+75, 6+25	96± 3	6	A+75, 6+85	158± 4*	10
A+75, 7+25	225± 7*	14	A+75, 7+75	328± 8*	22

Yalues exceeding background plus 2 standard deviations.

(2529n-11)

TABLE 2-2 (CONTINUED)

LARGE AREA GAMMA RAY SURVEY RESULTS

JANUARY 23 - FEBRUARY 8. 1990

Location	Counts Per Minute	uR/hr	Location	Counts Per Minute	uR/hr
A+75, 7+75	488±10*	35	A+75, 8+25	144± 5*	9
B+25, 1+25	116± 3	7	B+25, 1+75	106± 3	6
B+25, 2+25	118± 3	7	B+25, 2+75	118± 3	8
B+25, 3+25	125± 4	8	B+25, 3+50	127± 4	8
B+25, 4+25	130± 4	9	B+25, 4+75	143± 4	10
B+25, 5+25	113 <u>±</u> 3	8	B+25, 5+75	114± 3	8
B+25, 6+25	129± 4	9	B+25, 6+75	84± 3	6
B+25, 7+25	258± 7*	13	B+25, 7+75	298± 8*	14
B+25, 8+25	149 <u>+</u> 4*	9	B+75, 1+25	120± 3	6
B+75, 1+75	112± 3	7	B+75, 2+25	104± 3	7
B+75, 2+75	100± 3	7	B+75, 3+25	97± 3	7
B+75, 3+75	115± 3	8	B+75, 4+25	85± 3	6
B+75, 4+75	110± 3	9	B+75, 5+25	111± 3	7.5
B+75, 5+75	105 <u>+</u> 3	8	B+75, 6+25	120± 3	8
B+75, 6+75	157± 4*	11	B+75, 7+25	85± 3	7
B+75, 7+75	74± 3	7	C+25, 1+25	98± 3	7
C+25, 1+75	90± 3	: 6	C+25, 2+25	96± 3	8
C+25, 3+75	96± 3	7	C+25, 4+25	132± 4	9
C+25, 4+75	104± 3	7	C+25, 5+25	135± 4	8
C+25, 5+75	133 <u>+</u> 4	7.5	C+25, 6+25	99± 3	8
C+25, 6+75	117± 3	9	C+25, 7+25	128± 4	10

Values exceeding background plus 2 standard deviations.

(2529n-12)

TABLE 2-2 (CONTINUED)

LARGE AREA GAMMA RAY SURVEY RESULTS

JANUARY 23 - FEBRUARY 8, 1990

Location	Counts Per Minute	uR/hr	Location	Counts Per <u>Minute</u>	uR/hr
C+25, 7+75	111± 3	11	C+75, 1+25	102± 3	6.5
C+75, 1+75	103± 3	7	C+75, 2+25	99± 3	· 6
C+75, 2+75	89± 3	6	C+75, 3+25	106± 3	7
C+75, 3+75	136± 4	8	C+75, 4+25	81± 3	5
C+75, 4+75	80± 3	5	C+75, 5+25	81± 3	5.5
C+75, 5+75	135± 4	9.5	C+75, 6+25	116± 3	8
C+75, 6+75	132± 4	9.5	C+75, 7+25	155± 4*	10
C+75, 7+75	140± 4	9	C+75, 8+25	118± 3	7
D+25, 1+25	Water	-	D+25, 1+75	98± 3	6
D+25, 2+25	91± 3	6	D+25, 2+75	89± 3	6
D+25, 3+25	90± 3	7	D+25, 3+75	90± 3	6
D+25, 4+25	66± 3	6	D+25, 4+75	72± 3	5
D+25, 5+25	91± 3	5.5	D+25, 5+75	115± 3	7
D+25, 6+25	116± 3	8	D+25, 6+75	132± 4	8
D+25, 7+75	145± 4	8	D+25, 8+15	134± 4	9
D+75, 8+15	132± 4	, 9	D+75, 7+75	126 <u>+</u> 4	8.5
D+75, 7+25	96± 3	6.5	D+75, 6+75	93± 3	6.5
D+75, 6+25	108± 3	7.5	D+75, 5+75	89± 3	6.0
D+75, 4+75	98± 3	8.5	D+75, 4+25	88± 3	6
D+75, 5+25	82± 3	6	E+25, 4+25	84± 3	5
E+25, 4+75	85± 3	6	E+25, 5+25	97± 3	6

^{*} Values exceeding background plus 2 standard deviations. (2529n-13)

TABLE 2-2 (CONTINUED)

LARGE AREA GAMMA RAY SURVEY RESULTS

JANUARY 23 - FEBRUARY 8, 1990

Location	Counts Per <u>Minute</u>	uR/hr	Location	Counts Per <u>Minute</u>	uR/hr
E+25, 5+75	97± 3	6	E+25, 6+25	102± 3	7
E+25, 6+75	111± 3	8	E+25, 7+25	103± 3	8
E+25, 7+65	128± 4	9	E+75, 7+65	128± 4	9
E+75, 7+25	102± 4	7	E+75, 6+75	98± 4	7
E+75, 6+25	88± 4	7	E+75, 5+75	93± 4	6
E+75, 5+25	77± 3	6	E+75, 4+75	87± 4	6
E+75, 4+25	110± 4	7	F+25, 4+25	139± 4	6
F+25, 4+75	114± 4	6	F+25, 5+25	114 <u>±</u> 4	7
F+25, 5+75	103± 4	7	F+25, 6+25	112± 4	6
F+25, 6+75	135± 4	7	F+25, 7+25	151± 5*	9
F+25, 7+75	124 <u>±</u> 4	8	F+75, 7+75	116± 4	8
F+75, 7+25	109± 4	7	F+75, 6+75	80± 3	5
F+75, 6+25	96± 4	7	F+75, 5+75	105± 4	6
F+75, 5+25	89± 4	7	F+75, 4+75	86± 3	6
F+75, 4+25	107± 4	7	G+25, 4+25	90± 4	5
G+25, 4+75	95± 4	7	G+25, 5+25	99± 4	6
G+25, 5+75	87± 4	. 7	G+25, 6+25	84± 3	7
G+25, 6+75	72± 3	5	G+25, 7+50	44± 2	5
G+75, 7+70	68 <u>+</u> 3	5	G+75, 6+75	76± 3	5
G+75, 6+25	95± 4	6	G+75, 5+75	113± 4	8
G+75, 5+25	114± 4	7.5	G+75, 4+75	117± 4	7
G+75, 4+25	86± 4	6			

Values exceeding background plus 2 standard deviations.

(2529n-14)

TABLE 2-2 (CONTINUED)

LARGE AREA GAMMA RAY SURVEY RESULTS

JANUARY 23 - FEBRUARY 8. 1990

<u>Location</u>	Counts Per <u>Minute</u>	uR/hr	Location	Counts Per <u>Minute</u>	uR/hr
H+25, 7+70	65 <u>+</u> 3	5	H+25, 6+75	52± 3	5
H+25, 6+25	94± 4	6	H+25, 5+75	114± 4	8
H+25, 5+25	126± 4	8	H+25, 4+75	109 <u>±</u> 4	8.5
H+25, 4+25	105± 4	6	H+75, 7+70	66± 3	4.5
H+75, 6+75	66± 3	5	H+75, 5+75	93± 4	7
H+75, 5+25	114 <u>±</u> 4	7 .	H+75, 4+75	92± 4	7
H+75, 4+25	92± 4	6.5	H+75, 3+75	102± 4	6.5
I+25, 7+70	70± 3	4.5	I+25, 6+75	69± 3	5
I+25, 5+75	98± 4	7	I+25, 5+25	114± 4	7
I+25, 4+75	111± 4	7	I+25, 4+25	111± 4	8
I+25, 3+75	101± 4	6.5	I+75, 7+70	84± 3	7
I+75, 5+75	105± 4	6	I+75, 4+75	99± 4	7.5
I+75, 4+25	135± 4	8	I+75, 3+75	117± 4	9
I+75, 3+25	99± 4	8	J+25, 3+75	86± 3	7
J+25, 3+25	107± 4	8	J+25, 4+75	91± 4	6
J+25, 4+25	102± 4	7.5	J+75, 6+15	84± 3	6
J+25, 5+25	106± 4	7.5	J+25, 5+75	100± 4	7
J+25, 6+25	90± 4	6.5	J+25, 6+60	132± 4	8
J+75, 5+75	73± 3	4	J+75, 5+25	72± 2	4
J+75, 4+75	162± 5*	9	J+75, 4+25	85± 3	6
J+75, 3+75	74± 3	5			

^{*} Values exceeding background plus 2 standard deviations.

(2529n-15)

TABLE 2-2 (CONTINUED)

LARGE AREA GAMMA RAY SURVEY RESULTS

JANUARY 23 - FEBRUARY 8, 1990

Location	Counts Per Minute	uR/hr	Location	Counts Per Minute	uR/hr
M+25, 2+75	118± 4	8	M+25, 3+25	141± 8*	8
M+25, 3+75	142± 5*	7.5	M+25, 4+25	106± 4	5.5
M+25, 5+25	60± 3	4	M+25, Fence	66± 3	4
M+75, Fence	60± 3	4	M+75, 5+25	64± 3	4
M+75, 4+25	115± 4	6	M+75, 3+75	126± 4	6.5
M+75, 3+25	106± 4	9	M+75, 2+75	91± 4	9
M+75, 2+25	113± 4	7.5	M+75, 1+75	112± 4	7.5
N+25, 1+75	83± 3	8	N+25, 2+25	108± 4	7
N+25, 2+75	96± 4	10	N+25, 3+25	102± 4	7
N+25, 3+75	95± 4	6.5	N+25, 4+25	73± 3	5
N+25, 5+25	71± 3	5	N+25, Fence	68± 3	4
N+75, 4+75	70± 3	5	N+75, 4+25	103± 4	7.5
N+75, 3+75	88± 4	7.5	N+75, 3+25	114 <u>±</u> 4	8
N+75, 2+75	110± 4	8	N+75, 2+25	89± 4	9.5
N+75, 1+75	94± 4	9	N+75, Bulkhead	89± 4	8
O+25, Bulkhea	d 104± 4	7	0+25, 1+75	123± 4	7
0+25, 2+25	118± 4	7	0+25, 2+75	107± 4	8
0+25, 3+25	107± 4	. 8	0+25, 3+75	96± 4	7
0+25, 4+25	143± 5	9	0+75, 3+75	112± 4	7

^{*} Values exceeding background plus 2 standard deviations.

TABLE 2-2 (CONTINUED)

LARGE AREA GAMMA RAY SURVEY RESULTS

JANUARY 23 - FEBRUARY 8, 1990

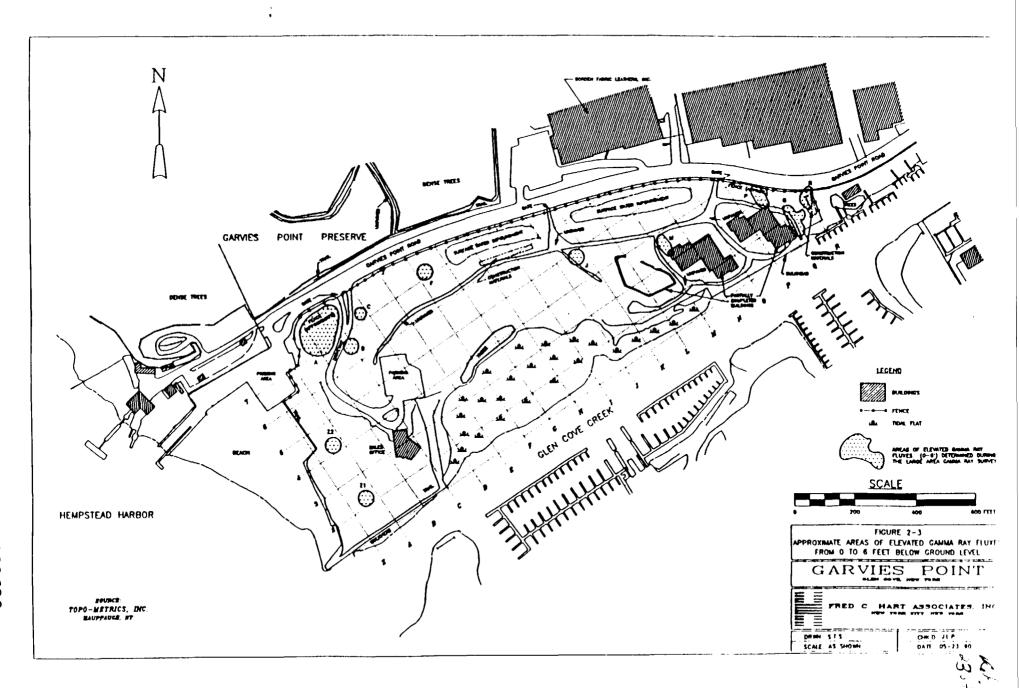
Location	Counts Per Minute	uR/hr	Location	Counts Per Minute	uR/hr
0+75, 3+25	118± 4	8	0+75, 2+75	62± 3	. 6
0+75, 2+25	137± 4	10	0+75, 1+75	131± 4	9
0+75, Bulkhead	107± 4	8	P+25, Bulkhead	129± 4	9.5
P+25, 1+75	133± 4	11	P+25, 2+25	114 <u>±</u> 4	11
P+25, 2+75	136± 4	11.5	P+25, 3+25	108± 4	9
P+25, 3+75	99± 4	7	P+75, 3+25	214± 6*	17
P+75, 2+75	190± 5*	20	P+75, 2+25	96± 4	8
P+75, 1+75	124 <u>±</u> 4	12	P+75, 1+25	82± 3	9
P+75, Bulkhead	87± 4	8	Q+25, 1+25	132± 4	9
Q+25, 1+50	2369± 34*	120	Q+25, 1+75	171± 9*	11.5
Q+25, 2+25	516± 16*	26	Q+25, 2+75	120 <u>±</u> 8	7.5
Q+25, 3+10	118± 8	6.5	Q+50, 1+75	312 <u>+</u> 12*	19
Q+75, 1+25	116± 8	5	Q+75, 1+75	130± 8	9
Q+75, 2+25	106± 7	6	R+O, 1+25	114± 8	4.5
R+O, 2+0	152± 9*	8	R+O, 2+25	138± 8*	8
R+15, 1+50	113± 8	6	R+25, 2+0	140± 8	6.5
R+25, 2+25	533± 16*	[:] 21			

^{*} Values exceeding background plus 2 standard deviations.

TABLE 2-3

LIST OF GRID POINTS THAT EXCEED BACKGROUND LEVELS

Grid Point	Location on Map in Figure 2-3
Z+75, 2+25	21
Z+75, 4+25	72
A+75, 6+85 A+75, 7+25	A A
A+75, 7+75	A
A+75, 8+25	Å
B+25, 7+25 B+25, 7+75	A
B+25, 7+75 B+25, 8+25	A A
<i>5</i> 723, 6723	^
B+75, 6+75	. В
C+75, 7+25	С
F+25, 7+25	F
J+75, 4+75	J
M+25, 3+25	н
M+25, 3+75	Ä
P+75, 3+25	
P+75, 2+25	P.
Q+25, 1+50	Q
Q+25, 1+75	Q Q
0+25, 2+25	Q
Q+50, 1+75	Q
R+10, 2+0	R
R+O, 2+25	R
R+25, 2+25	R



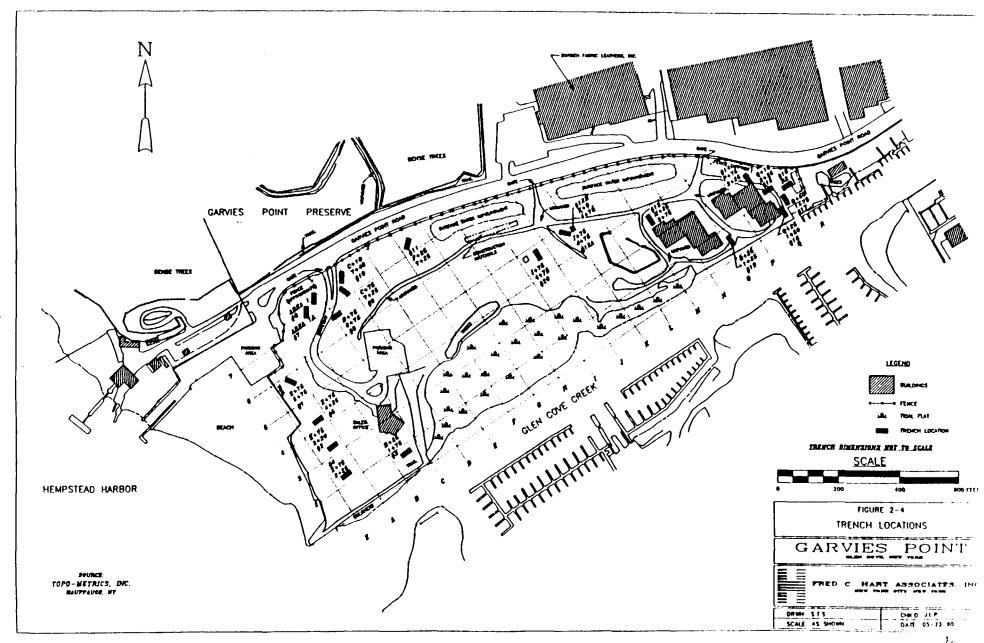


During the investigation, there was some concern that the results of the large area gamma ray detector survey could be affected by encountering soil densities of greater than 1.5 g/cc such as in areas where stone or concrete might have been dumped. An increase in soil density would reduce the effective depth of ore detection to less than 1 meter. Fortunately, concrete and stone were observed in only a few locations. Soil densities were estimated from the weights of the radionuclide samples collected and were found to range from 0.7 to 1.3. Thus, the estimated depth of detection of 1 to 2 meters was valid throughout the majority of the property. Actual ore layers were detected by the gamma spectrometer at depths of up to four feet at locations where microR meters showed only background levels.

2.4 Phase II Subsurface Investigation

- 2.4.1 Objective. The purpose of the subsurface investigation was to allow a visual inspection of the composition and depth of the fill material in areas of both elevated gamma ray fluxes and in background areas. Soil samples were collected for laboratory analysis to measure the range of radionuclide concentrations in the soils.
- 2.4.2 <u>Methodologies</u>. The trenching activities took place between February 9 and 13, 1990. All trenches were dug by Direct Environmental, Inc. of Hest Babylon, New York, using a JD 590 Trachoe and were overseen by HART and NDL personnel.

Fifteen trenches were dug in the areas of elevated gamma ray fluxes and an additional five trenches were dug at locations where background gamma ray fluxes were measured. The locations of the trenches are shown in Figure 2-4 and Table 2-4 shows the correlation between trench locations and the 25 areas of elevated gamma ray fluxes noted on Table 2-3. The trenches were approximately 3 feet wide by 5 to 15 feet in depth and up to 50 feet in length. Trench logs were filled out at each trench location by a HART geologist. Information noted on the logs included trench location and identification number, the start and finish dates, the condition and composition of the trench walls, the sample collection depths, air



27.70

TABLE 2-4

CORRELATION BETHEEN GAMMA RAY FLUX MEASUREMENTS AND TRENCH LOCATIONS

Grid Points at which at which Elevated Gamma Ray Fluxes Were Noted	Location	NDL Trench Number and Coordiates of Trenches Excavated for Subsurface Investigation
-	Background	Number 1, 2+75, 5+75
-	Background	Number 2, Z+75, 3+20
-	Background	Number 3, B+25, 1+25
Z+75, 2+25	Z 1	Number 4, Z+75, 2+25
Z+75, 4+25	22	Number 5, Z+75, 4+25
A+75, 6+85 A+75, 7+25 A+75, 7+75 A+75, 8+25 B+25, 7+75 B+25, 7+75 B+25, 8+25	A A A A A	Number 6, Area 1, B+10, 7+75 Number 7, Area 1A, A+75, 6+85
B+75, 6+75	В	Number 8, B+75, 6+75
C+75, 7+75	C	Number 9, C+75, 7+25 to 7+75 Number 10, C+10, 7+40
F+25, 7+25	F	Number 11, F+25, 7+25
J+75, 4+75	j	Number 12A, J+75, 4+75 Number 12B, K+0, 4+75
M+25, 3+25 M+25, 3+75	H	Number 13, M+25, 3+75
P+75, 3+25 P+75, 2+75	P P	Number 14, P+75, 3+25
Q+25, 2+25 Q+25, 1+50 Q+25, 1+75 Q+50, 1+75	Q : Q Q	Number 15, Q+10, 2+25 Number 16, Q+25, 1+50 Number 17, Q+50, 1+75
R+0, 2+10 R+10, 2+25 R+25, 2+25	R R R	Number 18, R2 (R+O, 2+O)
-	Background	Number 19, 0+25, 1+25
-	Background	Number 20, 1+75, 4+25
(2529n-23)		

monitoring readings, and the length and depth of each trench. The logs are contained in Appendix B.

During trenching activities, the work zone was monitored with a microR meter, a particulate dust monitor (PDM), a photoionization detector (PID or HnU unit) and a combustible gas indicator (CGI). Drager Tubes for vinyl chloride monitoring were available on-site in the event organic vapor concentrations exceeded three parts per million (ppm) for one minute. Five air monitoring stations were set up around the site to monitor airborne particulates in the vicinity of the work area. Two stations were set up upwind and three stations were situated downwind of the trenching activities. All work was performed in Level C protective gear in order to prevent contact with or inhalation of radionuclides in soil.

Samples were obtained by a HART geologist by collecting soil from the appropriate interval directly from the bucket of the trachoe. The instability of the trench walls made it impossible for HART personnel to enter the pits. The trachoe operator cleared away soil that fell into the trench from the sidewalls and collected a soil sample from an undisturbed location on the bottom of the trench. Soil samples were collected at two foot intervals in each trench using this technique. Each sample was properly identified, packed in coolers and documented under full chain-of-custody procedures. The samples were directly relinquished to the NDL health physicist. All samples were analyzed by the NDL Organization, Inc. of Peekskill, NY, which participates in the EPA-NV quality assurance program.

After each trench was examined, logged and sampled, the large area gamma ray detector was lowered into the trench if it was not readily apparent that elevated gamma ray fluxes were present. The detector was not lowered into any trenches known to have elevated gamma ray fluxes. By lowering the detector to the trench bottom at six to eight feet below ground level, gamma ray fluxes at depths between six feet and native soil could be measured. In this way, it was possible to evalute gamma ray fluxes through the entire thickness of the fill. The trenches were

backfilled with soil, to the best extent possible, in the order in which the soils were excavated. After backfilling, the filled areas were surveyed for any exposed ore. The trachoe was surveyed for contamination each day and underwent decontamination at the Li Tungsten site at the completion of trenching activities.

2.4.3 <u>Results</u>. A total of 66 soil samples from the trenches were analyzed on an intrinsic germanium detector with a computer-based multichannel analyzer by the NDL Organization, Inc. Spectral data was reduced to radionuclide concentrations by the use of a gamma ray spectrum analysis program, the Oak Ridge National Laboratory "FUEL" gamma library, and a National Bureau of Standards mixed gamma calibration source prepared in the same geometry as the Garvies Point samples. A summary of the sampling results is shown in Table 2-6. No samples were collected from the background trenches 1,3, or 20 or from trenches 5 and 10. No samples were collected from trench 10 because the fill material encountered was difficult to sample; the lack of samples from trench 5 was an oversight. A copy of NDL's sampling report is contained in Appendix C.

The concentrations of thorium generally ranged from below the detection limit to about 28.5 pCi/g. One anomalously high concentration of thorium of 583 pCi/g was detected near the driveway at a depth of four to six feet in Area 1. Anomalously high concentrations of uranium and Ra-226 were also found in this sample from Area 1 at four to six feet. In the remaining samples, uranium concentrations ranged from below detection to about 57.3 pCi/g, and Ra-226 readings ranged from below detection to about 54.5 pCi/g.

The four upwind and four downwind air samples were analyzed and showed no detectable levels of radioactivity ($<4 \times 10^{-13}$ uCi/ml).

29.670

TABLE 2-5

SUMMARY OF RADIONUCLIDE CONCENTRATIONS
IN TRENCH SAMPLES

NDL No.	Trench Location	Sample Depth (feet)	NDL Sample Number	Th-nat pC1/g	U-nat pC1/g	Ra-226 pCi/g
2.	Z+75, 3+20	0 2 4 6 8	10 64 7 56 55	1.1 0.9 <0.9 <0.8 <0.7	<3.9 <2.9 <2.9 <1.3 <2.3	1.1 0.9 0.5 0.4 0.3
4.	Z+75, Z+25	0 2 4 6 8	19 20 8 18 9	<1.4 0.6 0.6 <1.3 <0.6	<3.1 <2.1 <1.4 <1.9 <1.4	0.6 0.6 0.3 0.5 0.4
6.	Area 1 (B+10, 7+75)	0 2 4 6 10	13 58 54 4 65	0.9 0.6 583 1.1 4.1	<3.4 <2.9 662 52.8 <6.1	0.9 0.7 772 2.7 3.1
7.	Area 1A (A+75, 6+85)	0 2 4 6	52 16 1 11	2.6 3.3 28.5 0.5	<5.9 <6.3 49.7 <1.5	3.7 3.4 47.4 <0.3
8.	B+75, 6+75	0 2 4 6 8 12-16	61 5 57 14 62 59	0.7 <0.6 28.1 19.3 4.5 11.6	<3.0 <2.3 44.5 18.8 6.9 13.9	0.4 0.4 41.3 26.4 6.8 17.8
9.	C+75, 7+25 to 75	0 2 4 6 8	17 60 63 66 22	0.8 0.8 1.8 <1.1 3.0	<2.1 <2.0 <2.1 <2.0 <8.3	0.6 0.8 1.2 0.5 4.2
10.	C+10, 7+40	4	15	4.0	<10.9	6.6
11.	F+25, 7+25	0 2 6	23 3 2	<0.9 1.1 2.1	<1.5 <2.7 <1.9	0.4 0.6 1.1

(2529n-26)

TABLE 2-5

SUMMARY OF RADIONUCLIDE CONCENTRATIONS IN TEST PIT SAMPLES (CONTINUED)

NDL No.	Trench Location	Sample Depth (feet)	NDL Sample Number	Th-nat pCi/g	U-nat pCi/g	Ra-226 <u>pC1/g</u>
12A.	J+75, 4+75	0 2 6	6 53 21	0.4 0.8 0.7	<2.5 <1.4 <3.3	0.5 0.6 0.4
13.	M+25, 3+75	0 2 4 6	26 12 25 24	<0.8 0.8 <0.8 <0.6	<1.9 <1.8 <3.8 <2.5	<0.4 0.6 0.4 0.4
14.	P+75, 3+25	2 4 6 8 10	43 45 46 48 38	<0.7 0.6 12.4 0.6 0.9	<2.0 <1.8 20.8 <2.4 <3.0	0.8 0.7 16.9 0.7 0.6
15.	Area 3 (Q+10, 2+25)	0 2 4 6	39 44 42 51	24.4 <0.7 9.5 4.8	57.3 <4.6 13.7 <9.3	54.5 0.8 17.6 6.4
16.	Area 2 (Q+25, 1+50)	0 2 4 6 8	37 36 35 47 50	2.7 0.7 0.7 1.2 1.3	<5.3 <2.6 <2.8 <4.2 <4.8	0.9 <0.4 <0.4 0.5 1.9
17.	Q+50, 1+75	0 2 4 8 10	29 34 27 28 49	4.0 <1.0 4.3 1.7 0.5	<12.5 <3.9 <10.0 <6.5 <2.6	3.3 0.5 8.3 4.0
18.	R+O, 2+O (R stake)	0 2 6 10	30 33 32 41	0.6 3.9 2.9 <1.8	<1.8 <9.7 <6.6 <3.5	0.4 6.8 4.3 1.5
19.	0+25, 1+25	6 8	31 40	4.0 <1.0	<7.2 ·	11.1

1×0:10 30 of 70

TABLE 2-5

SUMMARY OF RADIONUCLIDE CONCENTRATIONS IN TEST PIT SAMPLES (CONTINUED)

NDL No.	Trench Location	Sample Depth (feet)	NDL Sample <u>Number</u>	Th-nat pCi/g	U-nat pCi/g	, Ra-226 pC1/g
12A.	J+75, 4+75	0 2 6	6 53 21	0.4 0.8 0.7	<2.5 <1.4 <3.3	0.5 0.6 0.4
13.	M+25, 3+75	0 2 4 6	26 12 25 24	<0.8 0.8 <0.8 <0.6	<1.9 <1.8 <3.8 <2.5	<0.4 0.6 0.4 0.4
14.	P+75, 3+25	2 4 6 8 10	43 45 46 48 38	<0.7 0.6 12.4 0.6 0.9	<2.0 <1.8 20.8 <2.4 <3.0	0.8 0.7 16.9 0.7 0.6
15.	Area 3 (Q+10, 2+25)	0 2 4 6	39 44 42 51	24.4 <0.7 9.5 4.8	57.3 <4.6 13.7 <9.3	54.5 0.8 17.6 6.4
16.	Area 2 (Q+25, 1+50)	0 2 4 6 8	37 36 35 47 50	2.7 0.7 0.7 1.2 1.3	<5.3 <2.6 <2.8 <4.2 <4.8	0.9 <0.4 <0.4 0.5 1.9
17.	Q+50, 1+75	0 2 4 8 10	29 34 27 28 49	4.0 <1.0 4.3 1.7 0.5	<12.5 <3.9 <10.0 <6.5 <2.6	3.3 0.5 8.3 4.0
18.	R+O, 2+O (R stake)	0 2 6 10	30 33 32 41	0.6 3.9 2.9 <1.8	<1.8 <9.7 <6.6 <3.5	0.4 6.8 4.3 1.5
19.	0+25, 1+25	6 8	31 40	4.0 <1.0	<7.2 ·	11.1

(2529n-27)

3.0 DISCUSSION

3.1 Description of Site Soils

Based on the results of the subsurface investigation, there is a significant layer of fill material overlying most of the site. The nature and content of the fill was fairly consistent throughout the site. The most common debris found in the fill material included plastic trash bags, wood, bricks, glass, metal, tires, concrete and paper. At locations Z+75, 2+75 and M+25, 3+75 what appeared to be bluish-purple paint and dye waste was observed. On the eastern side of the site near locations Q+25, 1+50; Q+50, 1+75 and Q+10, 2+25, the fill was comprised of sandy soil with minor amounts of debris. Native soil encountered at the site was a reddish-brown medium-grained sand with gravel.

The fill appeared to be the thinnest, as expected based on the aerial photograph review, near the northern and western boundaries of the property. The fill thickness on the western boundary at grid location Z+75, 5+75 was approximately 3.5 feet. On the northern boundary, at location F+25, 7+25, no fill was observed. The native soil was not observed in any of the trenches on the eastern portion of the site even though trenches of up to 10 feet in depth were excavated. Groundwater was encountered in several trenches in the central and western portions of the site at approximately 8 to 10 feet below the surface.

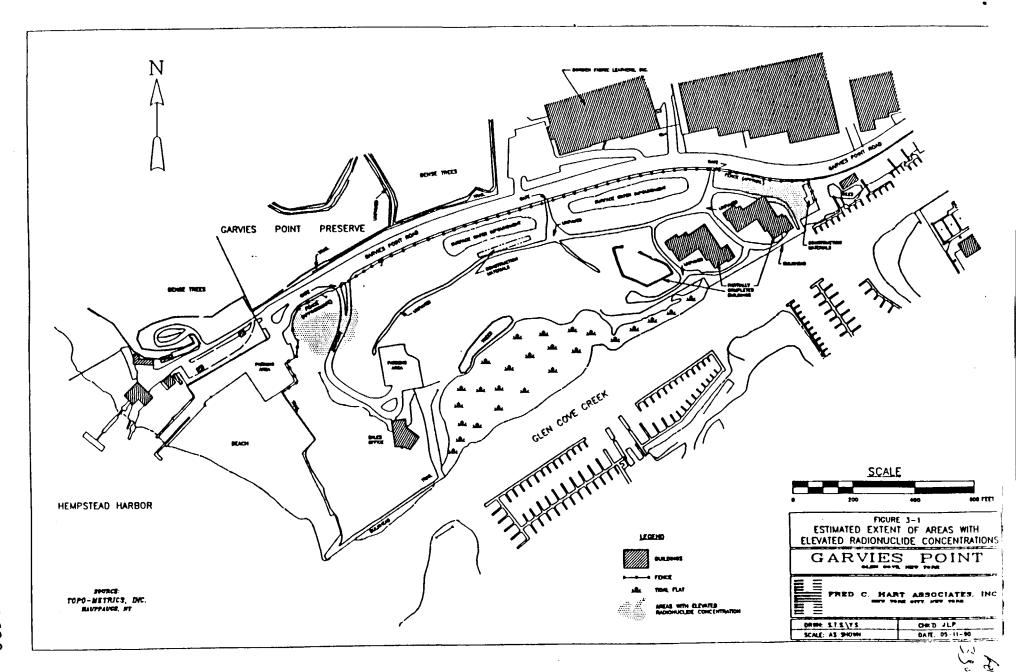
3.2 Soil Quality

During the large area gamma ray detector survey, elevated gamma ray fluxes were measured at 25 grid points. Most of these grid points were located in one of two general areas of the property: around the main entrance/driveway area, and in the far eastern corner of the property. These results correlate well with the data collected during the Phase I survey. The area in the driveway encompasses the original Area I from the Phase I investigation and the eastern area encompasses both locations on the eastern side of the site, designated as Areas 2 and 3, at which elevated readings were detected in the Phase I survey.

Six grid points outside of the areas designated as 1, 2 and 3 in Phase I showed elevated gamma ray fluxes during the large area gamma ray survey: Z1, Z2, F, J, M and Q+25, 1+50 (Figure 2-3). Trenches were dug at these locations to determine the source of the elevated readings. The sample results from these locations show acceptable radionuclide concentrations generally ranging from less than 1 pCi/g to 2 pCi/g. It is thought that the elevated gamma flux readings are from clean soil that was slightly enriched in thorium either as a result of natural processes or as a result of debris such as firebrick, lantern mantles, zircon process sands or small stray pieces of slag from Li Tungsten.

The sampling results from the trenches confirmed that the material containing elevated levels of radionuclides was generally found in two areas; one on the far eastern side of the property (Areas P. Q. and R) and one on the far western side near the entrance/driveway (Areas A. B. and C). It is interesting to note that the most elevated readings of thorium. uranium, and radium are found in a fairly discrete zone in both locations at about 4 to 8 feet below ground surface. In both of these areas, the elevated readings are associated with a black powder or granular material which contained concentrations of uranium and thorium series usually in the 1 to 50 pCi/g range. The physical appearance and corresponding radionuclide concentration range of this material closely matches that of the lower level tungsten ores found on Parcels A and B of the Li Tungsten Readings above the 4-6 foot layer tend to be close to background and in most cases, readings from below the 4-6 foot layer are close to background as well. Minor exceptions to this trend were found in trenches 8, 15 and 17. At trench 8, elevated concentrations of uranium and thorium were found at depths ranging from 4 to 16 feet. The elevated readings are directly associated with a black granular material in this trench. Slightly elevated readings in the 0-2 foot zone were found at two locations: trench 15 and trench 17 on the eastern side of the property.

An estimate of the areal extent of soils with elevated levels of radionuclides is shown in Figure 3-1. The area near the entranceway is approximately 28,750 square feet and the area in the eastern corner is



approximately 18,750 square feet. The thickness of the material containing the black granular ore varies from location to location within each area, but is an average of eight feet thick. Therefore, the total volume of soil in this area is approximately 14,000 cu yds.

3.3 Impact of Findings on Remedial Investigation

Based upon the results of both the large area gamma ray detector survey and the analysis of samples from the trenches. Ted Rahon, the NDL health physicist has determined that the Remedial Investigation can be conducted as planned with some minor modifications to the Health and Safety Plan. During any test boring activities within either of the two zones of elevated radionuclide concentrations, respirators and tyvek should be worn and all soil samples should be screened with a microR meter. Since the trenching activities did not generate any detectable levels of radionuclides in the air in the vicinity of the work area, it is unlikely that the boring activities, which cause less soil disturbance, would cause any particulate generation. Therefore, the only changes necessary to the Health and Safety plan are those mentioned above for on-site worker protection. The revised sections of the Health and Safety Plan are shown in Figure 3-2.

Access to the entire site has been restricted by the construction of a fence around all sides of the site not adjacent to the creek. In addition, "No Trespassing" signs have been posted. In order to provide protection to anyone gaining unauthorized access to the site, the areas in the vicinity of the driveway and on the eastern side of the site will be roped off and "Do Not Enter" signs will be posted. Since the majority of the site showed below background levels of radionuclides, no other access restrictions are necessary.

Figure 3-2

TABLE 8-1

FIELD INVESTIGATION TEAM

SITE SAFETY PLAN

A. GENERAL INFORMATION

SITE: Garvies Point

PROJECT NO.: 130032

LOCATION: Glen Cove, New York

PREPARED BY: John Persico

DATE: January 25, 1989

REVISED BY: Laura Truettner DATE: May 11, 1990

OBJECTIVE(S): Drill test borings, install groundwater monitoring wells and conduct air, subsurface soil and groundwater sampling to identify

soil and groundwater contamination, if any.

PROPOSED DATE(S) OF INVESTIGATION: Summer 1990

BACKGROUND REVIEW

PRELIMINARY:

COMPLETE: X

DOCUMENTATION/SUMMARY: OVERALL HAZARD: SERIOUS MODERATE LOW X UNKNOWN

B. SITE/MATERIAL CHARACTERISTICS

MATERIAL TYPE(S): LIQUID

SOLID X

SLUDGE X

GAS X

CHARACTERISTIC(S): CORROSIVE

IGNITABLE

RADIOACTIVE

VOLATILE X

TOXIC X REACTIVE

UNKNOWN

OTHER (NAME):

SITE DESCRIPTION: The site has been used as a disposal area for sediment dredged from Glen Cove Creek, low level ore from Li Tungsten and as a municipal landfill.

PRINCIPAL DISPOSAL METHOD (type and location): Surface dumping.

STATUS (active, inactive, unknown): Inactive

HISTORY: Sediment dredged from Glen Cove Creek by the Army Corps of Engineers was disposed of at the site in 1960 and 1965. From 1971 to the early 1980s, the site was used as a municipal landfill. Incinerator ash, wastewater treatment plant sludge, and household and street debris were deposited on the site during this period. Soil samples collected in 1985 contained metals and estimated levels of pesticides and PCBs and a groundwater sample from one upgradient well contained several volatile compounds in concentrations exceeding Class GA guidelines. Soil samples collected in 1989 and 1990 showed elevated levels of radionuclides in two discrete areas of the site (Figure 8-1).

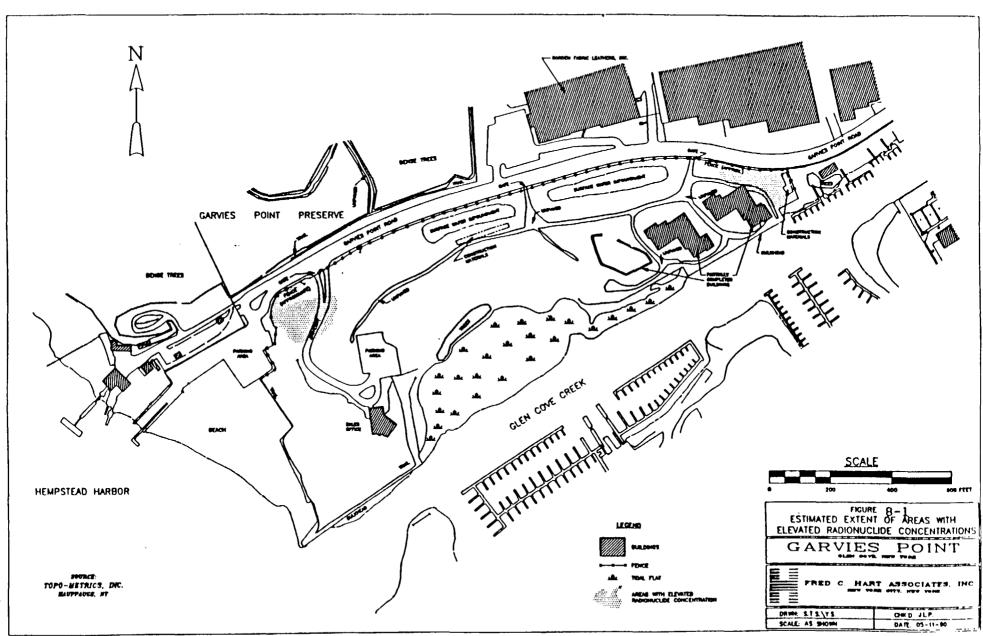


Figure 3-2 (Continued)

TABLE 8-1 (CONTINUED) C. HAZARD EVALUATION

Based on the 1985 and 1990 samples, a low potential exists for exposure at the site. Some particulate matter may be dispersed into the air during soil disturbance activities and a particulate dust monitor should be used to monitor these concentrations. In addition, when working in the areas shown in Figure 8-1, respirators and tyvek must be worn and all soil samples and sampling equipment should be screened with a microR meter. Certain hazards typically associated with landfills, such as production of methane or other organic vapors, must also be addressed.

D. SITE SAFETY WORK PLAN

PERIMETER ESTABLISHMENT: MAP/SKETCH ATTACHED? Yes SITE SECURED? Yes

PERIMETER IDENTIFIED? Yes ZONE(S) OF CONTAMINATION IDENTIFIED? Yes, radiological zones of contamination are identified

PERSONNEL PROTECTION:

LEVEL OF PROTECTION: D (with Level C equipment available on-site for all work conducted in radiological exclusion zones).

SURVEILLANCE EQUIPMENT AND MATERIALS: Photoionization detector, Drager air monitoring kit with tubes for vinyl chloride, combustible gas indicator, and personal particulate dust monitors, microR meter.

DECONTAMINATION PROCEDURES: All sampling equipment will be decontaminated between each use with the following procedure: detergent and water wash, distilled water rinse, nitric acid rinse, acetone or methanol rinse, hexane rinse, air dry. One decontamination area will be established on-site for steam cleaning and a separate station for hand and boot washing and disposal of personnel equipment will be established. Personnel equipment will be held on-site until sampling results become available, at which time it will be appropriately disposed of.

SPECIAL EQUIPMENT, FACILITIES, OR PROCEDURES: None

PREMISES ENTRY PROCEDURES: To be arranged with Village Green Realty at Garvies Point, Inc.

TEAM MEMBER (Major)

RESPONSIBILITY

James Perazzo
Laura Truettner
Peter Conde
John Persico
WORK LIMITATIONS (time of day, etc.):
Project Director
Project Manager
Field Team Leader
Site Safety Officer
Daylight hours

INVESTIGATION-DERIVED MATERIAL DISPOSAL: All development and purge water will be collected in 55 gallon drums and placed in an on-site, lined pool for temporary storage. A sample will be collected from the pool and (2567n-34)

Figure 3-2 (Continued)

TABLE 8-1 (CONTINUED)

analyzed for TCL organics and TAL inorganics to determine appropriate on-site or off-site disposal alternatives. Drill cuttings will be left on-site at each boring location.

E. EMERGENCY INFORMATION

LOCAL RESOURCES

Nassau County Police 911 POLICE:

City of Glen Cove Police (516) 676-1000

FIRE DEPARTMENT: City of Glen Cove Fire Dept. (516) 676-0366

EXPLOSIVES UNIT: City of Glen Cove Police (516) 676-1000

AMBULANCE: City of Gien Cove Police (516) 676-1000 (request ambulance)

HOSPITAL EMERGENCY ROOM: Glen Cove Community Hospital (516) 676-5000

(request emergency room)

POISON CONTROL CENTER: Nassau County Medical Center, Uniondale

(516) 542-2323

SITE RESOURCES

WATER SUPPLY: To be arranged with Village Green Realty at Garvies Point.

Inc.

TELEPHONE: To be arranged with Village Green Realty at Garvies Point.

Inc.

RADIO: n/a

OTHER: n/a

EMERGENCY CONTACTS

POSITION	<u>PERSONNEL</u>	PHONE		
CORPORATE SAFETY DIRECTOR	Larry Kaufman	(609) 663-0440		
PROJECT DIRECTOR	James Perazzo	(212) 840-3990		
CLIENT CONTACT	Eric Zoellner	(301) 727-3351		
NYSDEC CONTACT	Christopher Magee	(518) 457-5637		

(2567n-35)

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Figure 3-2 (Continued)
TABLE 8-1 (CONTINUED)

F. EMERGENCY ROUTES

Directions to Glen Cove Community Hospital: Take Garvies Point Road east. Make right onto Herb Hill Road, proceed to Charles Street, make right turn and continue to traffic light. Make left onto Forest Avenue. Take Forest Avenue north approximately 1 mile to Walnut Road. Right turn onto Walnut Road, take Walnut Road 1 block to St. Andrews Lane. Make right turn onto St. Andrews Lane and left to emergency room (see Figure 8-2 for map).

Appendix A

Calculation of Effective Depth of Ore Detection

CALCULATION OF EFFECTIVE DEPTH OF ORE DETECTION

Assumptions:

Source: 5 meter diameter x 0.5 meter thick slab of

thorium-bearing material (Thorium Conc: 100 pCi/g)

Source material density: 3 g/cc; self-absorp. factor: 0.17

Gamma fraction for T1-208 2.6 MeV from Th-232 parent assuming chain equilibrium: 36%

Detector area: 324 sq.cm.

Soil densities = 1.0, 1.2 and 1.5 g/cc

BKG on 1 foot on sand: 40 cpm

Efficiency of 8"x2" CsI crystal (photofraction + single escape peak for 2.6 MeV - determined experimentally): 60%

Source strength:

100 pCi | 9.8 x 106 cc |0.17|3 g |2.22 dpm| 0.36 phot| 0.6 cts |
g | | | | cc | pCi | dis | photon |

= 2.4 x 108 cpm excluding geometry and overlaying soil attenuation

Depth (m)	Geometryl Factor		Atten. Factor	Count Rate above BKG (cpm)
1	5.3 x 10 ⁻⁴	1.5	2.5 x 10-3	315
		1.25	6.7×10^{-3}	855
		1.0	1.8 x 10-2	2250
1.5	4.0 x 10 ⁻⁴	1.5	1.2 x 10 ⁻⁴	12
		1.25	5.5 x 10-4	53
		1.0	2.5 x 10-3	240
2	3.0 x 10 ⁻⁴	1.5	6.1 x 10-6	0.4
		1.25	4.5 x 10-5	3.2
		1.0	6.8 x 10 ⁻⁵	4.9

1. from Principles of Nuclear Radiation Detection

Appendix Bottom Trench Logs



N ^-5.	-
--------	---

NDL No. 1

Trench No.

Z+75, 5+75

Project Name/Number

Garvies Point/ NY 102-01

Location

GlenCove

HART Inspector

Suzanne Morrissey

Excavator/Equip Operator Direct / Brent Thompson

Start/Finish Date 2 /9 /90 - 2/9/90

(0855) (1530)

Elevation: Ground

Condition of Trench

walls stable

Depth of Trench

8

REMARKS:

. selected as a" Clear" trench

· perchal tribling water

· all monitoring equipment reads back ground

Sketch of Trench Wall

ourface	Black-brown fill, brick fragments, medium-grained sand with gravel, auto parts, root fragments, metal shards, paper (Edecomposed)
31/4	
	grey silty sand grading to a reddish-brown silty Sand

54'

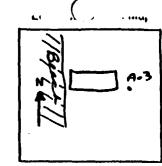
Brown gravelly sand, medium - coarse grained

4-5-4-

LENGTH _3

L03650





(5)

NDL No. 2

Trench No.

2+75,3+20

Project Name/Number

Garvies Point/Ny102-01

Location

Glen Cove

HART Inspector

Suzanne Morrissey

Excavator/Equip Operator Direct / Brent Thompson

Start/Finish Date

2/12/90 - 2/12/90 (0830) (1120)

Flevation: Ground

Condition of Trench

Stable walls

Depth of Trench

No groundwater

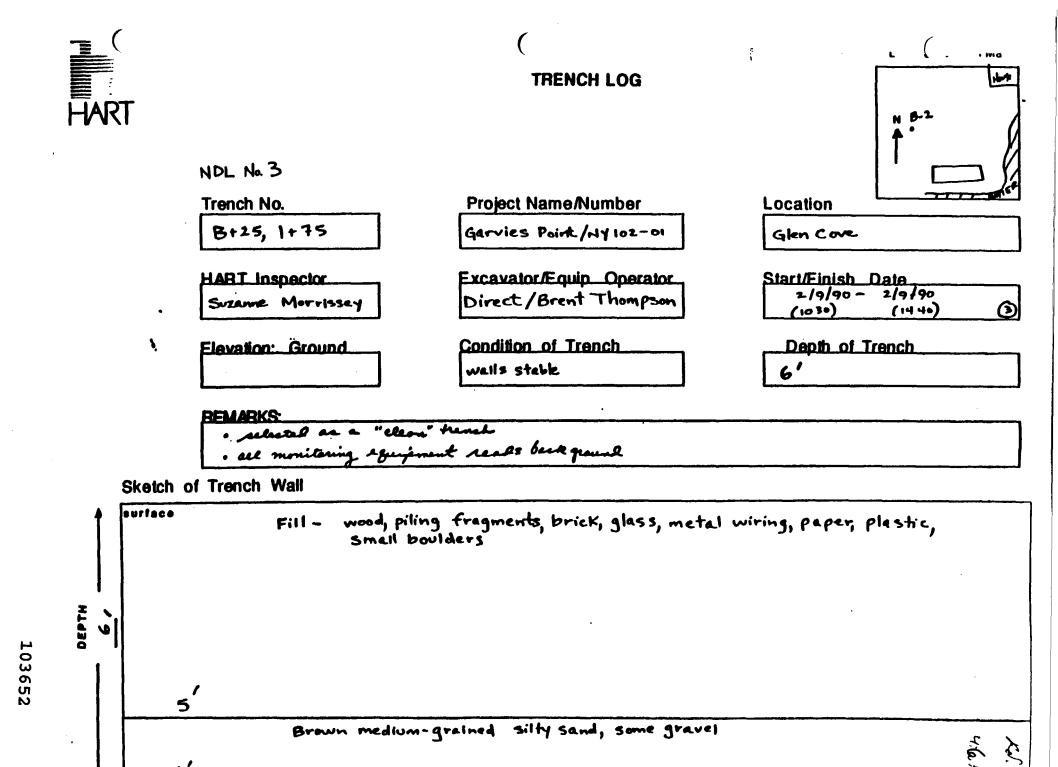
8'

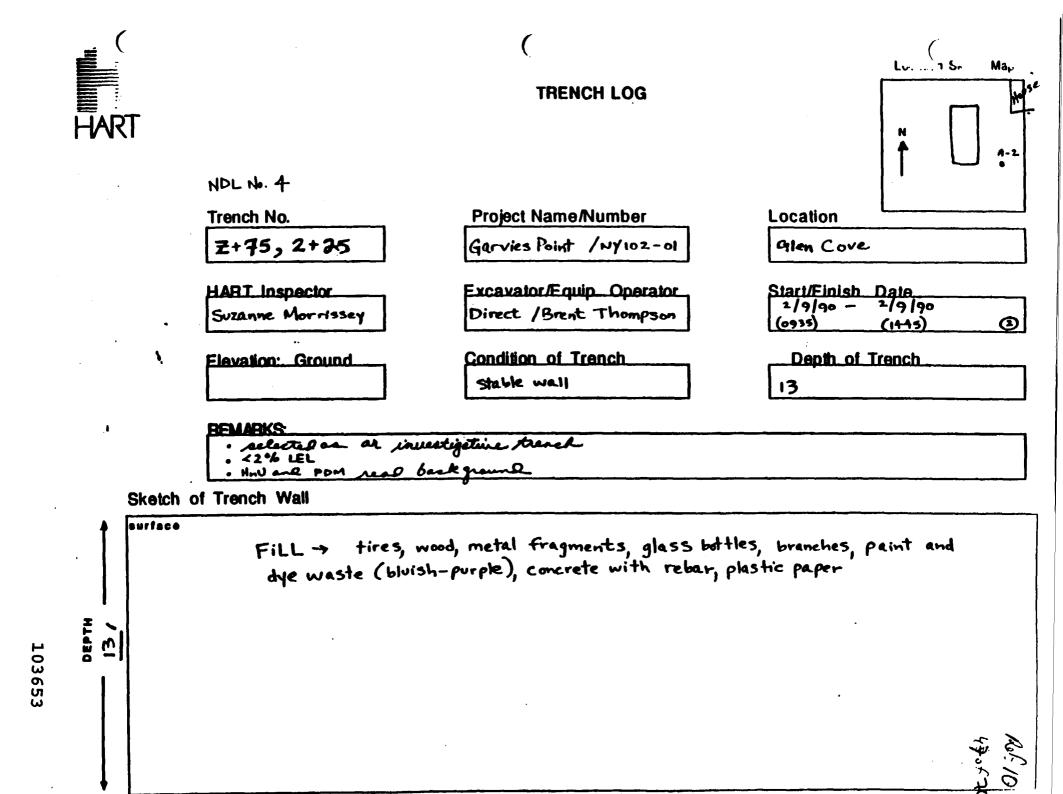
en collected at 0-6, 2,4,6,8, selected as a background hence ("there")
writering agriphent sends background except for I clay "chunk" (sed a gray)
is read ~ 11 uR/br.

Sketch of Trench Wall

leurtace Black fill, clay clumps (see remarks above), branches, concrete, tires, scrap metal, brick, newspaper Reddish-brown sand with coarse gravel

LENGTH 48







-		****
		#use
N		
	L	_ لـ

NDL No. 5

Trench No.

Z+75, 4+25

Project Name/Number

Garvies Point/HY102-01

Location

Glen Cove

HART Inspector

Suzanne Morrissey

Excavator/Equip Operator Direct /Brent Thompson

Start/Einish Data 2/9/90 - 2/9/90 (1115) (1435)

Elevation: Ground

Condition of Trench
walls colleged

Depth of Trench

10'

EIHVAIICH....GIUUHA

REMARKS:

. very damp at bottom but no groundwater

. pemples collected at 0-6, 2, 4,6, 8'

Sketch of Trench Wall

surface

FILL -> clothing, glass bottles, small boulders, metal fragments (some very large)

* Due to trench collapse, we had to dig a few feet deeper to place V-ray detector. It was visible that the fill extends to at least 12' deep.

10)

10365

DEPTH

LENGTH 41



Rose Rose

NDL No. 6

Trench No.

AREA 1

Project Name/Number

Garvies Point /HY 102-01

Location

Glen Cove

HART Inspector

Suzanne Morrissey

Excavator/Equip Operator Direct / Brent Thompson

Start/Finish Data 2/12/90 - 2/12/90

 $\frac{2}{12}\frac{90 - \frac{2}{12}90}{90}$

Elevation: Ground

Condition of Trench

no collapse

Depth of Trench

10'

REMARKS

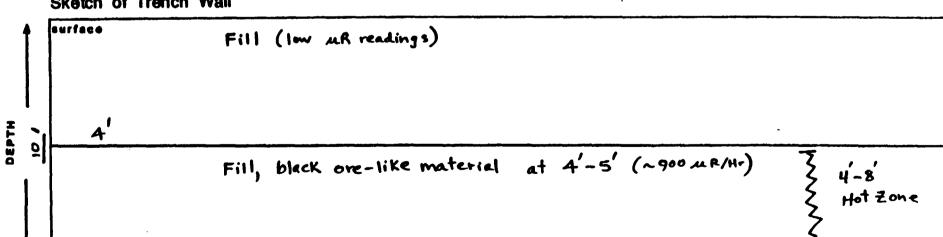
. samples collected at 0-6,2,4,6,10'

7 4' sample read 900-1,000 MR/hr in backhoe bucket

. Small trench, Ied Rahon didn't want to disturb too much "hot" fail

Sketch of Trench Wall

10



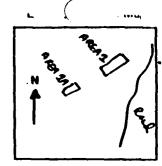
- LENGTH 10



trench at

Area 1

TRENCH LOG



(7)

NDL No. 7

Trench No.

AREA 1A

HART Inspector Suzanne Morrissey

Elevation: Ground

Project Name/Number

Garvies Point /NY102-01

Excavator/Equip Operator Direct / Brent Thompson

Condition of Trench small trench, stable wells Location

Glencove

Start/Finish_Date 2/12/90 - 2/12/90 (1045)

Depth of Trench

· pamples collected at 0-6", 2', 4', 6'

Sketch of Trench Wall

eurtace

Fill -> branches, plastic gloves, McDonald's cartons, plastic bags

41 -> ~ 90 MR/HR

5'- refusel, broke through a concrete layer

6 -> background readings, tan medium-grained sand with grave



3,4	
† 0	9
+i'dge	

(8)

NDL No. 8

Trench No.

B+75,6+75

Project Name/Number

Garries Point / NY102-01

Location

Glen cove

HART Inspector

Suzanne Morrissey

Excavator/Equip Operator Direct / Brent Thompson

Start/Finish Date

(1130) (1220)

Elevation: Ground

Condition of Trench

Depth of Trench

14'

REMARKS:

· pemples exclused at a-6, 2, 4,6,8, 12'

. offset ~18' se beause bookste operate feet shak in his binger while digging here

Sketch of Trench Wall

ourlace, 0-3' Brown Clean (?nobricks, etc.) fill soil

4' -> black fill (slag? fragments), ~ 100 MR/HR

Black fill, tree branches, pilings, tar-like substance, large boulders, ~ 30 MR/HR

+ |

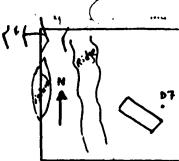
12

Groundwater M2-14'

LENGTH 15

7 1 1 1





9

NDL No. 9

Trench No.

C+75, 7+25

Project Name/Number

Gervies Point /Ny102-01

Location

Glen Cove

HART Inspector

Suzanne Morrissey

Excavator/Equip Operator Direct / Brent Thompson

Start/Einish Date

1/12/90- 2/12/90 1428) (1500)

Elevation: Ground

Condition of Trench

walls stable

Depth of Trench

8'

REMARKS

. pemples collected at 0-6,2,4,6,8'

· all monitaring equipment background (except at 8')

Sketch of Trench Wall

Black Fill -> glass, brick, pilings, cans, branches, large tree stump

8' - 28 MR/HR (8- 1463 counts/5 mins)

DEPT

103658

LENGTH 30



N A	
1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	
	.c4

(10)

NDL No. 10

Trench No.

C+10, 7+40

Project Name/Number

Garvies Point /NY 102-01

Location

Glen Cove

HART Inspector

Suzanne Morrissey

Excavator/Equip. Operator. Direct / Brent Thompson

Start/Finish Date

2/12/90 2/12/90-(1500)

(1510)

Elevation: Ground

Condition of Trench

walls stable

Depth of Trench

semple collected from 4' interval

Sketch of Trench Wall

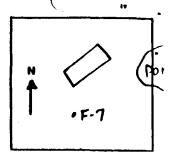
eurtace

Fill- brick, wood, logs or piling fragments, large boulders

2' -> 25 MR/M

DEPTH





NDL No. 11 Trench No.

F+25,7+25

Project Name/Number

Garvies Point /WY102-01

Location

Glen Cove

HART_Inspector

Suzanne Morrissey

Excavator/Equip_Operator_ Direct / Brent Thompson Start/Finish Date | 2/12/90

(1530) (1550)

Elevation: Ground

Condition of Trench walls very unstable, sand collepsing Depth of Trench

Sketch of Trench Wall

surface Brown, medium-grained sand, some gravel (No fill)

Black coarse sand, trace clay



L.	(Ma
N		• K5
†		

①

NDL No. 12 A

Trench No.

J+75,4+75

Project Name/Number

Garries Point/Ny102-01

Location

Glen Cove

HART Inspector

Suzanne Morrissey

Excavator/Equip Operator

Direct / Brent Thompson

Start/Einish Date

 $\frac{2}{12} \frac{40 - \frac{2}{12}}{90}$

Elevation: Ground

Condition of Tranch
Collapsing walls
(wet and full of garbage)

Depth of Trench

10

. Samples collected at 0-6", 2', 6'

Sketch of Trench Wall

eurface

•

Black fill - tires, wood and piling fragments, tank (?), glass bottles, film-like plastic (is possibly shredded trash bags), cloth, scrap metal

6' -> 470 LEL on CGI

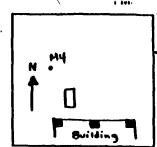
10' - 10% LEL . CGI

? Possible Groundwater at 10' (could be garbage waste)

£5.70 0.150

LENGTH 35





NDL No. 13

Trench No.

M+25, 3+75

Project Name/Number

Garvies Point/NY10Z-01

Location

Glen Cove

HART inspector

Suzanne Morrissey

Excavator/Equip Operator
Direct/Brent Thompson

Start/Finish Date

2/13/90 - 2/13/90 (0900) (0945)

(3)

Elevation: Ground

Condition of Trench

unstable met walls, no collapse

Depth of Trench

8'

REMARKS

· all monitoring equipment background

Sampler collected at 0-6"

Sketch of Trench Wall

surface

١

Brown fill - scrap metal, rubber, paint and dye waste

0-6" -> 10 ur/h

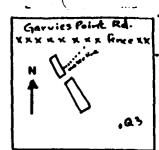
4-8' -> 7 me/m

B ,

Groundwater at 8'

2.5





NDL No. 14 Trench No.

P+75, 3+25

Project Name/Number

Garvies Point /NY102-01

Location

Glen Cove

HART Inspector

Suzanne Morrissey

Excavator/Equip. Operator Direct / Brent Thompson Start/Finish Date 2/13/90 - 2/13/90 (0945) (1040)

Elevation: Ground

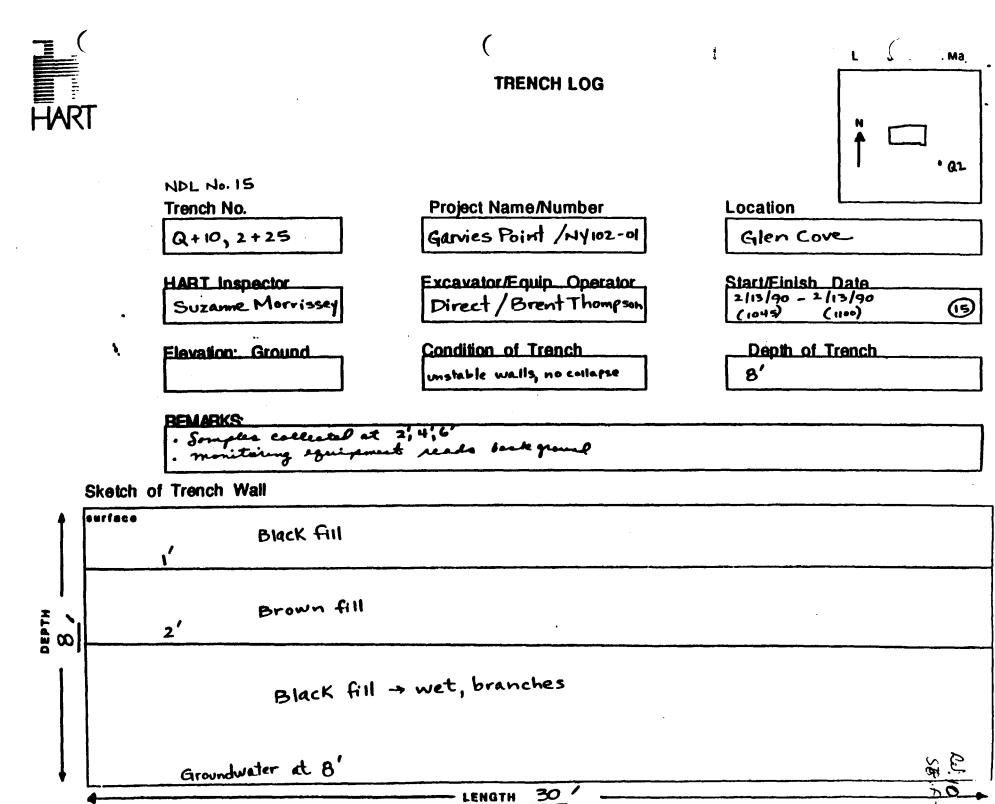
Condition of Trench walls unstable, partial cave-in Depth of Trench

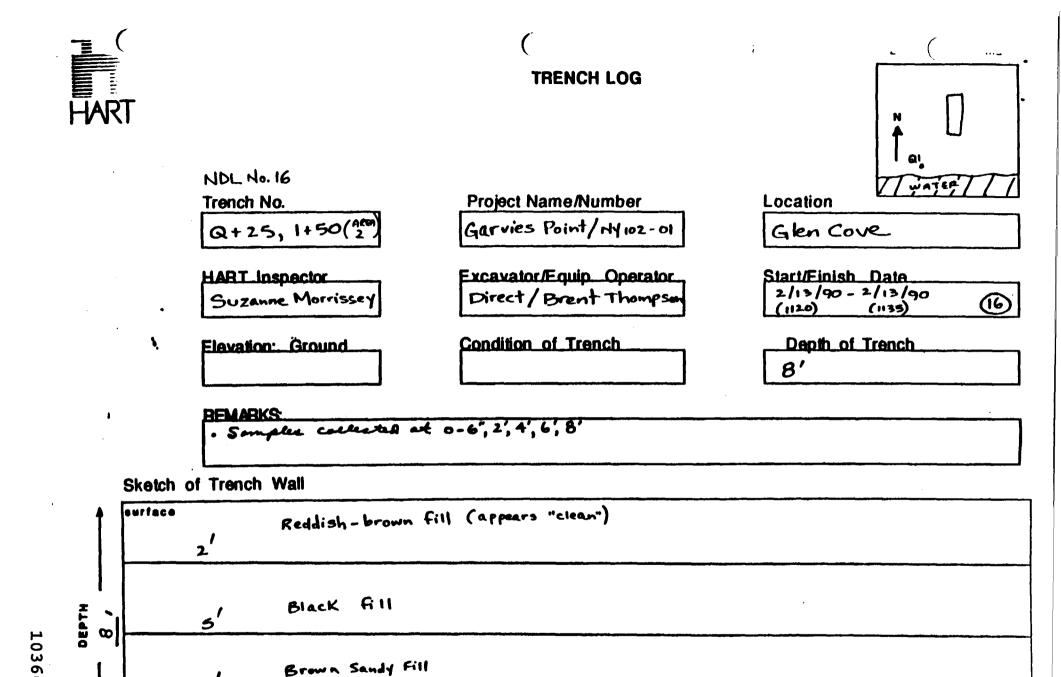
10'

Sketch of Trench Wall

surface Black and Brown fill - plastic bags, large tree stump

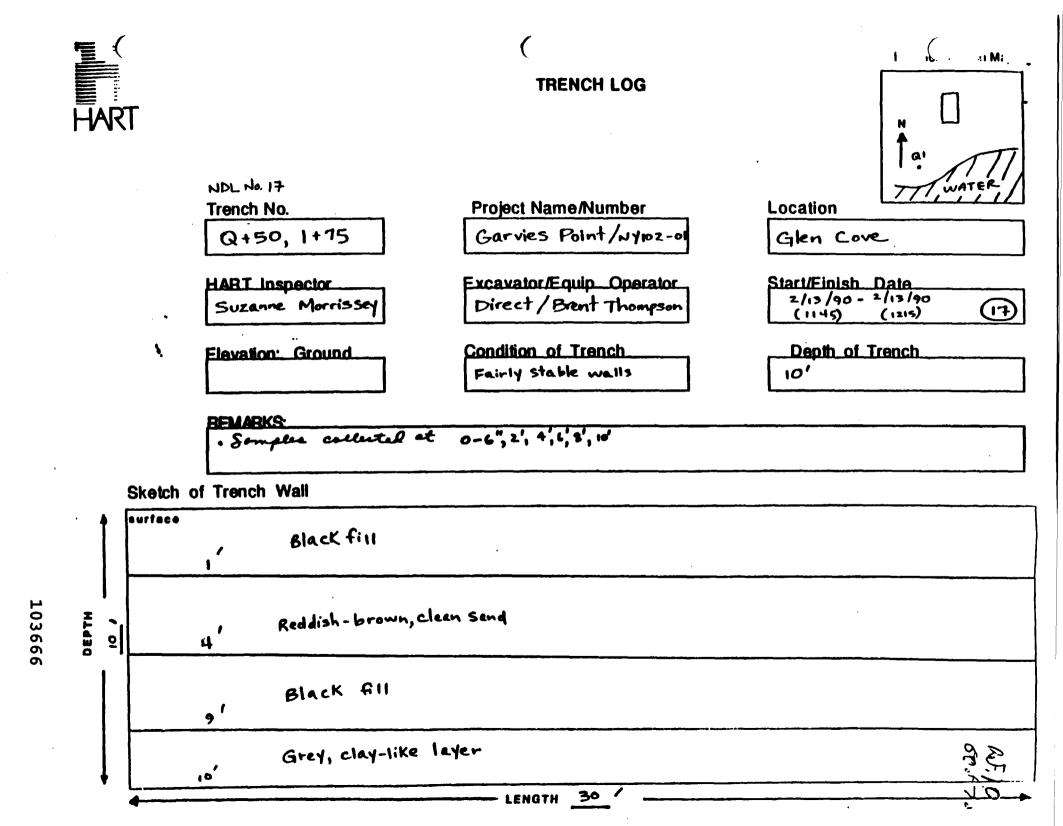
DEPTH





Black All, gravelly, somebrick

Grandwater at 8'





_ L		<i>y</i>	J	IVI G
N ↑	ſ		□ .	
11	7	A	TER	77/

NOL No. 18

Trench No.

RZ Stake

Project Name/Number

Garvies Point/Nyloz-01

Location

Glen Cove

HART Inspector

SUZANNE Morrissey

Excavator/Equip Operator

Direct / Brent Thompson

Start/Finish Date

2/13/90 - 2/13/90 (1400) (1345)

1

Elevation: Ground

Condition of Trench

walls semi-stable

Depth of Trench

Aamples collected at 0-6", 2', 6', 10'

Sketch of Trench Wall

surface Brown sendy fill Black fill - branches, brick, glass, metal shards

~8' - black, oily looking zone, oil-like odor and Sheen, shell + brick fragments



	6/43.
N (1)	اا
//water	77/

NDL No. 19

Trench No.

0+25, 1+25

Project Name/Number

Garvies Point/Hy102-01

Location

Glen cove

HART Inspector

Suzanne Morrissey

Excavator/Equip. Operator
Direct/Brent Thompson

Start/Finish Date 2/13/90 - 2/13/90 (1445) (1515)

(9)

Elevation: Ground

Condition of Trench

fairly stable

Depth of Trench

8'

REMARKS:

· Sompler coelested at B'

monitoring equipment reads back ground

Sketch of Trench Wall

surface

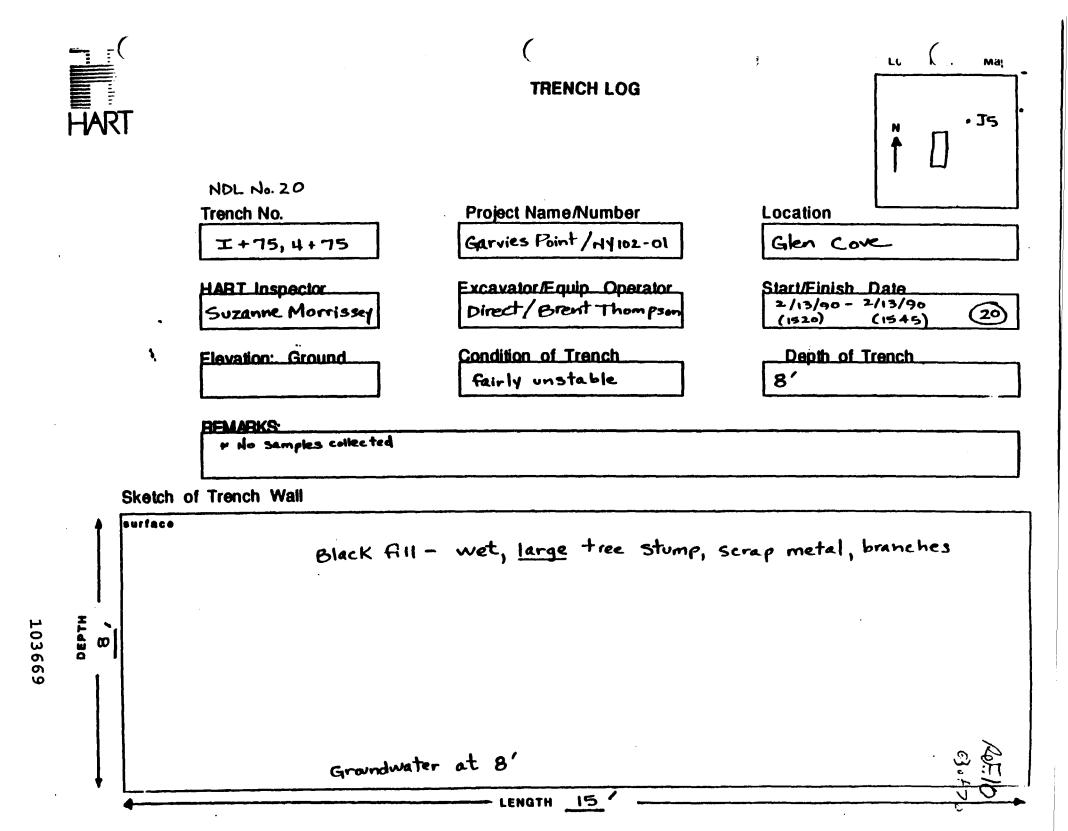
DEPTH 8

103668

Brownish - grey fill -> branches, scrap metal, brick, glass

LENGTH 40

65.70



Appendix C
Laboratory Analytical Data

		2.6	•		Soi!	uR/hr	MDL	Radionuc	lide Concer	ntration
Pit	Grid	MeV	Type of			€ 1cm		Th-nat	U-nat	Ra-225
#	Location	CPM	Trench*	(ft)	tion	fr soi!	1	(pCi/g)	(pCi/g)	(pCi/g)
1	z+75, 5+75	106	EKG	0	top scil	7-9				
	•			2	gray soil -	•				
	(W side of property)			4	some refuse gray soi! -	•				•
	footer of			-	some refuse				-	
				6	gray soi! -	•				
	1:	17.135	.181	8-10	some refuse yellow clay	•				
		f, mid	•							
2	Z+75 3+20	112	EKG	C	top soil	6-9	10	1.1+0.4	< 3.9	1.140.1
	400 1 1 2			2	gray soil -	•	64	0.9 <u>+</u> 0.4	< 2.9	0.9 <u>+</u> 0.1
	(W side of property)	154		4	some refuse gray soil -	*	7	< 0.9	< 2.9	0.5 <u>+</u> 0.1
	\$10\$01077	(S)			some refuse					_
				6	gray soil - some refuse	•	56	< 0.8	< 1.3	0.4 <u>+</u> 0.1
		180,	176	8	yellow clay	•	55	< C.7	< 2.3	0.3 <u>+</u> 0.1
		(H, :	mid)		•					
3	B+25, 1+75	106	EKG	0	top sail	6-8				
	/m =: 3			2	gray soil -	•				
	(W side of property)			4	some refuse gray soil -	•				
					some refuse					
		6,135 , mid		6	gray soil	**				
	3e	104	•		A		••			
4	Z+75, 2+25	194	Invest.	0	top soil	6-12	19	< 1.4	< 3.1	0.5 <u>+</u> 0.1
	(W side of			2	gray soil - some refuse	6-9	20	0.6 <u>+</u> 0.4	< 2.1	0.6 <u>+</u> 0.1
	property)			· 4	· •	•	8	0.6 <u>+</u> 0.3	< 1.4	0.3 <u>+</u> 0.1
				6	•	•	18	< 1.3	< 1.9	0.5 <u>+</u> 0.1
		150,10		8	•	•	9	< 0.6	< 1.4	0.4 <u>+</u> 0.2
		(E, mi	ra)	10	**	•				
		164 (F	7)	12	•	•				

*BKG - Background
Invest. - to investigate elevated.
veading

•		2.€				12/hr	NDL	Radionuclide Concentrat		tration
Pit	Grid Location	HeV CPH	Type of Trench*		tion	ê lem êr soi!		Th-mat (pCi/g)	U-nat (pCi/g)	Ra-226 (pCi/g)
5	Z+75,	208	Invest.	0	top soi!	8-10				
	4+25			2	gray soi! -	*				
	(W side of property)			4	some refuse	•				
		165	(H)	6	•	•				
				8	•	•				
		113, (mid	_	10	4	•				
6	Area 1 B+O,	488	Invest.	0	top soil	35	13	0.9 <u>+</u> 0.4	< 3.4	0.9 <u>+</u> 0.1
	7+75			2	brown soil black powder	- 900	58	0.6 <u>+</u> 0.4	< 2.9	0.7 <u>+</u> 0.1
	(W side of			4	a Pigor house:	*	54	583 <u>+2</u>	662 <u>+</u> 49	772 <u>+2</u>
	property)			6	• .	-	4	1.1 <u>+</u> 0.6	52.8 <u>+</u> 2.4	2.7 <u>+</u> 0.1
				8	*	•				
				10	brown sand	•	65	4.1 <u>+</u> 0.3	< 6.1	3.1 <u>+</u> 0.1
_										
7	Area la A+75,	158	Invest.	0	top soil	10	52	2.6 <u>+</u> 0.5	< 5.9	3.7 <u>+</u> 0.1
	6+85			2	brown soil	-	16	3.3 <u>+</u> 0.4	< 6.3	3.4 <u>+</u> 0.1
	(W side of property)			4	very hard, black layer	90	1	28.5 <u>+</u> 0.6	49.7 <u>+</u> 3.1	47.4 <u>+</u> 0.4
	·			6	brown sand	10	11	0.5 <u>+</u> 0.4	<1.5	<0.3

RUF.10

,										
Pit #	Grid Location	2.6 MeV CPH	Type of Trench*	Depth (ft)		uR/hr @ lcm fr soil	NDL Samp.	Radionuc! Th-nat (pCi/g)	ide Concer U-nat (pCi/g)	Ra-226 (pCi/g)
8	B+75, 6+75	157	Invest.	0	top soi! (on berm)	11	61	0.7 <u>+</u> 0.5	< 3.0	0.4 <u>+</u> 0.1
				2	brown soil	•	5	< 0.6	< 2.3	0.4 <u>+</u> 0.2
•	(W side - drwway berm)			4	black granula:	30-40	57	28.1 <u>+</u> 0.7	44.5 <u>+</u> 3.0	41.3 <u>+</u> 0.4
				6	•	•	14	19.3 <u>+</u> 0.5	18.8±2.1	26.4 <u>+</u> 0.3
•				8		•	62	4.5 <u>+</u> 0.3	6.9 <u>+</u> 1.6	6.8 <u>+</u> 0.2
				10		•				
				12-16	н	•	59	11.6 <u>+</u> 0.4	13.9 <u>+</u> 2.1	17.8 <u>+</u> 0.2
9	C+75, 7+25 to	155	Invest.	0	top soil	10	17	0.8 <u>+</u> 0.5	< 2.1	0.6 <u>+</u> 0.1
	7+75			2 .	prown/Aejjom (ou perm)	20	60	0.8 <u>+</u> 0.5	< 2.0	0.8 <u>+</u> 0.1
	(W side - dryway be:	m)		4	clay	35	63	1.8 <u>+</u> 0.4	< 2.1	1.2 <u>+</u> 0.1
				6	black clay-like	20	66	< 1.1	< 2.0	0.5 <u>+</u> 0.1
		292		8	e in its	28	22	3.0 <u>+0</u> .4	< 8.3	4.2 <u>+</u> 0.2
10	C+10.	173	Invest.	0	top soil	10				
	7+40			1	(edge of drive bricks & rubb)					
	(W side - at dryway			2	brown soil &	25				
	pavement)			4	rocks					
				4	black powder	25	15	4.0 <u>+</u> 0.4	< 10.9	6.6 <u>+</u> 0.2
11	F+25,	151	Invest.	0	brown sand	10	23	< 0.9	< 1.5	0.4 <u>+</u> 0.1
	7+25			2	•	10	3	1.1 <u>+</u> 0.3	< 2.7	0.6 <u>+</u> 0.1
	(mid- property)	235		6	; ·	10	2	2.1+0.2	< 1.9	1.1+0.1
•				-			-			

R.10 G7.170

r sj			2.6			Soil :	na/hr	NDL	Radionuclide Concentration		
	Pit	Grid Location	KeV	Type of Trench*	Depth (ft)		e lon fr soi!	Samp.	Th-mat (pCi/g)	U-nat (pCi/g)	Ra-126 (pC1/g)
12	12a	J+75, 4+75	162	Invest.	0	soil/gravel	9	6	0.4 <u>+</u> 0.3	< 2.5	0.5 <u>+</u> 0.1
					ż	decaying	-	53	0.8 <u>+</u> 0.3	< 1.4	0.6 <u>+</u> 0.1
		(mid- property)			6	refuse "	•	21	0.7 <u>+</u> 0.5	< 3.3	0.4 <u>+</u> 0.1
	ь	K+0,	149	Invest.	0	soil/gravel	10				
		4+75			2	decaying	8-10				
					6	refuse "	8-10				
	13	M+25,	142	Invest.	0	gravel,	8	26	< 0.8	< 1.9	< 0.4
		3+75			2	waste concrete soil, refuse	•	12	0.8 <u>+</u> 0.5	< 1.8	0.6 <u>+</u> 0.1
		(near condo)			4	•	•	25	< 0.8	< 3.8	0.4 <u>+</u> 0.1
			143		6	ø	•	24	< 0.6	< 2.5	0.4 <u>+</u> 0.1
						÷					
	14	P÷75, 3+25	214	Invest.	0	brown soi! (near prop. fo	17 mce)	-			
		(E side			2	brown soil	14	43	< 0.7	< 2.0	0.8 <u>+</u> 0.1
		of prop.)			4	•	•	45	0.6±0.4	< 1.8	0.7 <u>+</u> 0.1
					6	black vein (2' thick)	30	46	12.4 <u>+</u> 0.4	20.8 <u>+</u> 2.4	16.9 <u>+</u> 0.2
					8	yellow/black mix	-	48	0.6 <u>+</u> 0.5	< 2.4	0.7 <u>+</u> 0.2
					10	gray sand/clay	10	38	0.9 <u>+</u> 0.4	< 3.0	0.6 <u>+</u> 0.1
					>10	light brown sam	nd -				

RE,10 68.70

3 4	2.6			Soil - ul	nR/hr	NDL	Radionuclide Concentration			
Pit #	Grid Location		Type of Trench*	Depth (ft)	Descrip-	e lon fr soi!	Samp.	Th-nat (pCi/g)	U-nat	Ra-226 (pCi/g)
15	Q+10, 2+25	-	Invest	. 0	brown soil (on dirt road		39 io)	24.4 <u>+</u> 0.8	57.3 <u>+</u> 4.3	54.5 <u>+</u> 0.5
	(Area 3)			0.5	black material (thin layer)	80				
	(E side of prop.)			2	brown/orange	15	44	< 0.7	< 4.6	0.8 <u>+</u> 0.1
	or brob.)			4	black material	75	42	9.5 <u>+</u> 0.4	13.7 <u>+</u> 2.0	17.6 <u>+</u> 0.2
				6		*	51	4.8 <u>+</u> 0.3	< 9.3	6.4 <u>+</u> 0.1
				8	gray sand/clay (water table)	10				
				>9	refuse	•				
16	Q+25,	2369	Invest.	. 0	yellow clay w/	120	37	2.7 <u>+</u> 0.3	< 5.3	0.9 <u>+</u> 0.1
	1+50 (Area 2)			2	black+bricks	14	36	0.7 <u>+</u> 0.4	< 2.6	< 0.4
	(E side of prop.)			4	light brown	20	35	0.7 <u>+</u> 0.4	< 2.8	< 0.4
	or prop.,			6	gray soil	15	47	1.2 <u>+</u> 0.4	< 4.2	0.5 <u>+</u> 0.1
				8	gray sand/clay (water table)	10	50	1.3 <u>+</u> 0.2	< 4.8	1.9 <u>+</u> 0.1
17	Q+50,	312	Invest.	•	gray soil	20	29	4 040 3	. 19 E	2 2 4 2
••	1+75	744	INVEST.					4.0±0.3		3.3 <u>+</u> 0.1
	(E side			2	light brown sand		34	< 1.0	< 3.9	0.5 <u>+</u> 0.1
	of prop.)			4	black material w/red bricks	25	27	4.3 <u>+</u> 0.4	< 10.0	8.3 <u>+</u> 0.2
				6	•	20				
				8	•	30	28	1.7 <u>+</u> 0.5	< 6.5	4.0 <u>+</u> 0.2
				10	gray mud (water table)	15	49	0.5 <u>+0</u> .4	< 2.6	1.0 <u>+</u> 0.2

> ;		2.6			Soi!	uR/hr	NDL	Radionuc	lide Conce	ntration
Pit	Grid Location	MeV CPM	Type of Trench*			e lcm fr soil	Samp.	Th-nat (pCi/g)	U-nat (pCi/g)	Ra-226 (pCi/g)
18	R+0, 2+5	152	Invest	. 0	brown soil	10	30	0.6 <u>+</u> 0.5	< 1.8	0.4 <u>+</u> 0.2
	(E side			2	•	20 (60 in 3	33	3.9 <u>+</u> 0.4	< 9.7	6.8 <u>+</u> 0.2
	of prop.)			4	black, yellow, gray mixed la	20	,			
				6	# # # # # # # # # # # # # # # # # # #	•	32	2.9 <u>+</u> 0.5	< 6.6	14.3 <u>+</u> 0.2
				8	oily clay	10				
				10	light gray cla (water table		41	< 1.8	< 3.5	1.5 <u>+</u> 0.1
19	0+25, 1+25	132	BKG	0	brown soil	10				
	(between			2	brown/crange sand	10				
	condos)			4		10				
		217		6	gray/black material	10	31	4.0 <u>+</u> 0.4	< 7.2 ·	11.1 <u>+</u> 0.2
				8	•	10	40	< 1.0	< 1.9	0.7 <u>+</u> 0.1
20	1+75, 4+25	135	BKG	0	gray soil					
	(mid-			2	garbage & ashes	8				
	property)			4	*	•				
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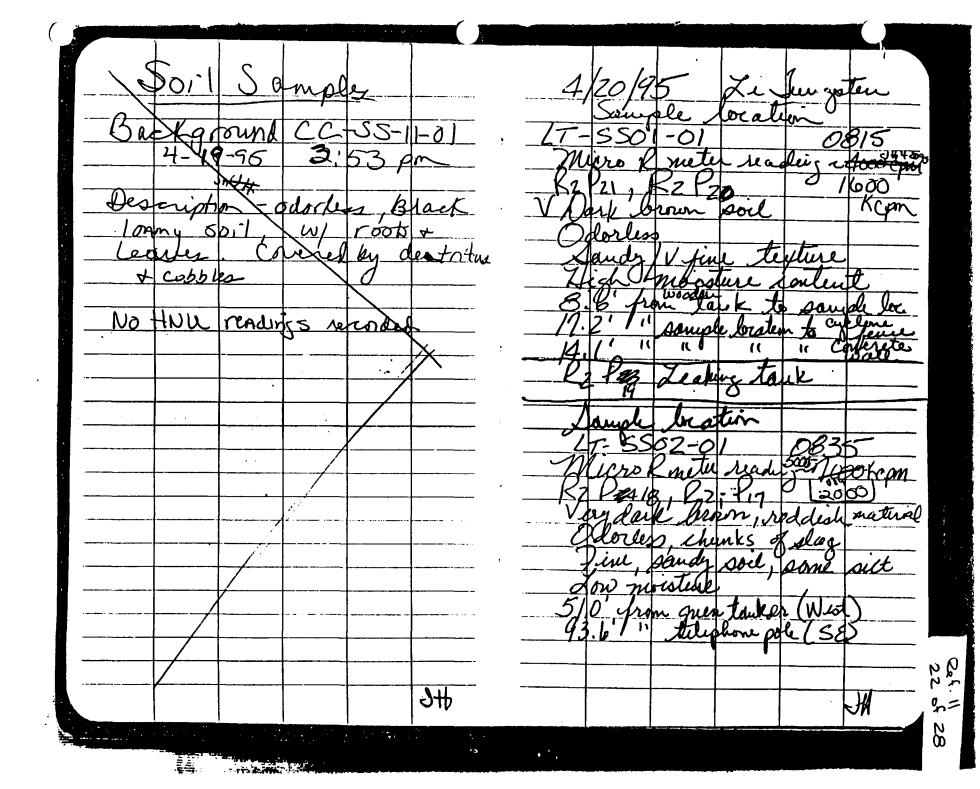
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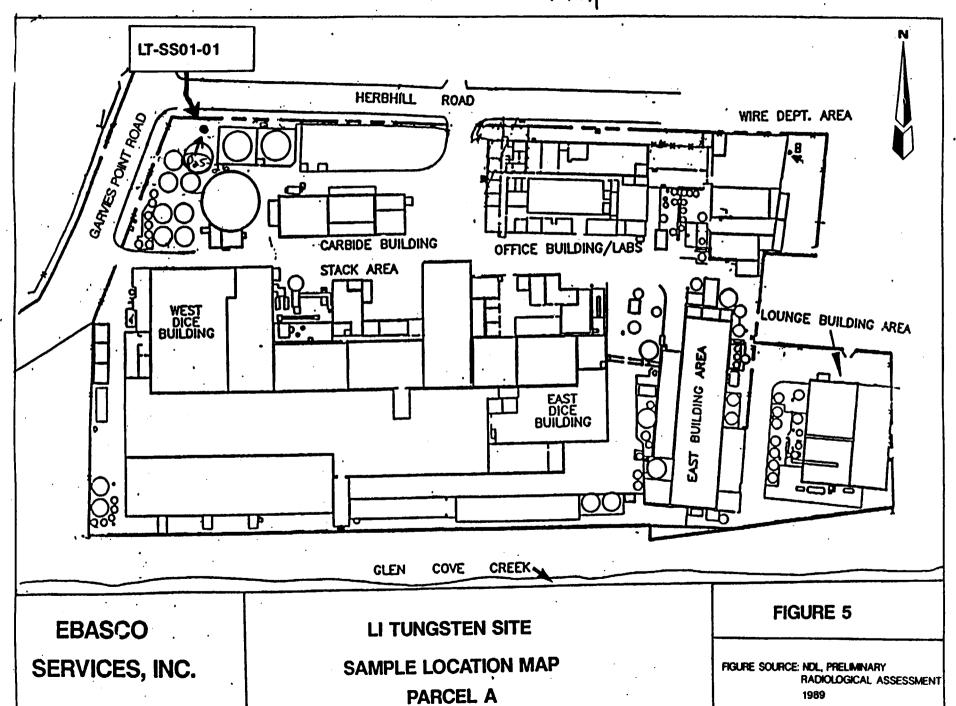
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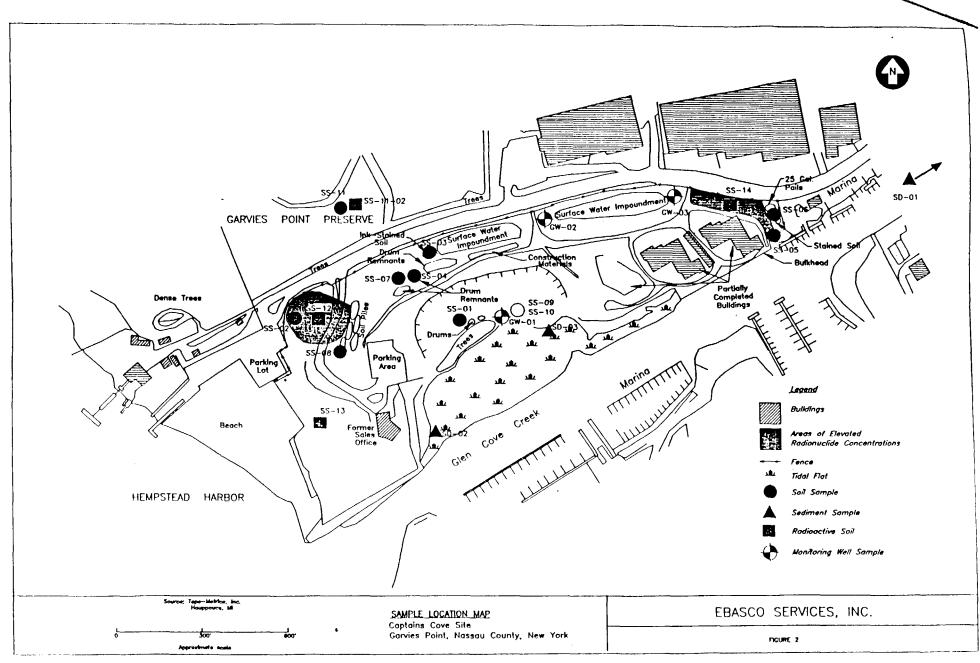
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Photo Location Map



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REFERENCE 12

SURFACE AND SUBSURFACE SOILS OBTAINED AT CAPTAIN'S COVE CONDOMINIUM SITE (ALL RESULTS IN MG/KG)

			CAPTAIN'S SURFACE S			CAPTAIN'S COVE BSURFACE SOILS	
CONTAMINANT	CC-SS11-02 (BACKGROUND TAKEN AT 0-6")	CC-SS11-02 (BACKGROUND TAKEN AT 4')	CC-SS14-01	CC-SS15-01 (DUPLICATE OF CC-SS14-01)	CC-SS12-01	CC-SS13-01	
TUNGSTEN	185 J	1.4 J	3,200 J	3,820 J	51	1,210 J	

SURFACE SOIL OBTAINED AT LI TUNGSTEN SITE (ALL RESULTS IN MG/KG)

	CC-SS11-62 (BACKGROUND TAKEN AT 0-6")		LT-SS02-01	LT-SS03-01	LT-SS04-01	LT-SS05-01	LT-SS05-01D
TUNGSTEN	185 J	3,050 J	16,200 J	1,160 J	4,540 J	1,160	1,420 J

20f34

CAPTAIN'S COVE
DATA VALIDATION REPORT
TUNGSTEN

Prepared by:

Cecelia N. Minch

Date:

2/1/95

BRIDGEPORT RENTAL AND OIL SERVICES DATA VALIDATION REPORT

SUMMARY:

This case consisted of 2 aqueous field blanks and 12 soil samples collected on April 20, 1995 and designated for the analysis of tungsten by ICP-MS method 200.8. One field duplicate pair (LTSSO5-01/05D) was collected and analyzed with satisfactory results. All soil results were reported on a dry weight basis. The % solid reported for sample CC-SS11-02 in this package was greater than 50%. However, the data user should be aware that in the report drafted for the metals analysis performed by IEANJ, the % solid was less than 50%. No action was taken.

Although a CLP package format was requested, several of the usual QC analyses were not performed since they are not specified in the method. In addition, since tungsten (W) is not listed on the target analyte list of the method, no CRDL was defined.

All data, however, were evaluated for Level IV DQO, employing USEPA Region II validation criteria. The specifics for each parameter and associated QC are detailed below.

The sample identifications used in this report have been truncated for expediency. Unless otherwise indicated, all sample IDs are suffixed with -Ol.

PRESERVATION:

The chains of custody indicated that the aqueous samples were preserved. The lab performed a check of the pH upon receipt, but did not provide documentation of the actual pH. Contact with the lab confirmed that all pH values were <2. No action was taken.

HOLDING TIMES:

All samples were prepped and analyzed within specified holding times.

MATRIX SPIKES:

The soil spike failed recovery criteria, but no action was required since the sample concentration was greater than 4 times the amount of spike added.

The aqueous spike was acceptable.

LAB DUPLICATES:

The % solids reported for the sample and lab duplicate varied by more than 1 %. Therefore, the reviewer converted the sample results to wet weight and recalculated the RPD, which met criteria. The aqueous duplicate was acceptable.

FIELD DUPLICATE:

The results of the field duplicate were acceptable.

LAB CONTROL SAMPLE (LCS):

The lab attempted to analyze a LCS, but no certified stock was available containing tungsten. The LCS that was analyzed did not contain any tungsten, so the results were not reported. No action was taken based on this criteria.

40F34

SERIAL DILUTION:

No serial dilution was performed. The following soil data were qualified as estimated (J) because the sample result exceeded 10 times the quantitation limit.

\$811-02, \$813, \$814, \$815, **\$801**, \$802, \$803, \$804, \$805-01D

No action was taken to the aqueous data since a serial dilution is not required to be performed on a field blank.

BLANK CONTAMINATION:

No qualifications were required.

INSTRUMENT CALIBRATION:

A CRI standard was not analyzed for W. No action was taken to the data since there was no specified CRDL.

INTERFERENCE CHECK SAMPLE (ICS):

An ICS was not performed. No interference was expected from the usual elements since the mass of interest for W is so high. No action was taken.

GENERAL COMMENTS:

Sample results were adjusted by the reviewer to correct for premature rounding performed by the laboratory.

The lab did not perform an IDL study or perform a linear range analysis. All samples were diluted to fall within the calibration range established by the initial calibration.

A from was not provided which summarized the results for the calibration blanks.

The client identifications for samples 9504608-08A and 09A required correction on the cross-reference supplied by the North Carolina lab.

The reported results for the soil spike required correction by the reviewer.

TELEPHONE RECORD LOG

	Date of Call:	6/21/95	
	Laboratory Name:	IEA	
•	Lab Contact:	Leanne	
	Client:	Foster Wheeler En	<u> </u>
	Client Contact:	C. Minch	
	Call Initiated By:	: Laboratory _	x Client
In :	reference to data for	or the following sample	number(s):
Captain's Cove ICP-MS data for Tungsten			
Sum	mary of Questions/Is	ssues Discussed:	
1. 7	CLP package format	was requested. On the	IEA COC, it was
indicated that analyses were to be performed per ILMO3.0 (CLP)			
protocol. Regardless of the method utilized, all of the usual			
OC should have been run. Please submit the following raw data			
8	nd/or summary form.	.	
	.) Percent solids de	eterminations. Were same	ole results reported
	on a dry weight b	casis?	
2	tuning solution a	malysis.	
3) CCB summary of re	sults.	
4) LCSS summary of r	results.	
5) IDL with date of	last determination.	
) linear ranges.		
2. F	lease submit a cros	ss-reference of sample 1	Ds with IEANC.
3. W	hy were prep blanks	diluted?	
4.1	n the method, the f	inal volume for soil pr	ceparations is
50 ml, but the runlog indicates 100ml. Was this taken into			
account during quantification?			
	Signat	ha M. Minch	6/31/95 Date
		·	

- 5. I cannot reproduce the reported results. Please supply a sample calculation for each matrix. Were results corrected for any interferences or blank subtracted? Include all necessary information to reproduce all values.
- 6. Why wasn't a serial dilution, interference check sample or CRI standard analyzed? They are CLP protocol for metals.
- 7. The wrong units were used for all aqueous data. Please resubmit.

6/21	IEANJ - Leanne: Briefly discuss items on faxed phone log.
6/22	IEANJ - Leanne: Informed me that requests were forwarded
·,	to IEANC and that she will be out of the office on
	6/23.
6/23	IEANJ: Mike left a message that resubs will be delayed.
6/26	IEANJ: Leanne called to say that the resubs should be
• –	faxed to her in the afternoon and fedex to me for
	6/27 AM.
6/27	IEANJ: Leanne called to say that the resubs are
-,	incomplete. Do I want a messenger to delivery a
	partial resub or wait for complete delivery on 6/28?
	I will wait for complete set.
6/28	IEANC: Message from G. Folk. An attempt to return the
0, 20	call was made at 5:01, but switchboard was off.
6/29	IEANC: Spoke with Gary regarding the unresolved
0/23	questions. He will convey my concerns to the
	inorganic manager and get back to me in PM. No
	return call.
6/30	IEANC: Gary was unable to effectively explain the
6/30	response to the dilution issues. I asked to speak
	with the IO manager D. Stogner. Spoke at length with
	Don regarding the prep and analysis. He will submit
	a brief explanation of the procedure and IDL
	determination.
7/1	IEANC: Spoke with don regarding the missing LCSS. He
	explained that no certified stock was available.
	Furthermore, the LCSS run appeared not to contain
- 4-	any W.
7/1	IEANJ: Spoke with Leanne to confirm that the pH was
	checked for the aqueous samples.

Memo

To:

Cecelia Minch

From:

Donald Stogner DC5

Subject:

Tungsten by Method 200.8

Date:

June 30, 1995

Please find listed below information to clarify how IEA performs 200.8 and information pertaining to your tungsten analysis specifically. Additionally I have included a copy of the dry weight log to aid in your calculations. If any questions are not answered here please do not hesitate to call.

IEA-NC performs 200.8 for soils by digesting one gram of sample using ultrex II grade acids following the steps listed in the 4.4 version of 200.8 from publication PB91-231498 section 11.2.2. After the digestion is complete IEA takes the sample to 100 ml and allows it to settle overnight. The method states to dilute the sample five fold prior to analysis. IEA performs this step immediately before analysis by pipetting 2 ml to 10 ml and adding internal standards. Should any reanalysis be required IEA repeats the dilution step from the one hundred ml final digested at either the required five fold dilution or higher. If the internal standards are outside the method specified range the lab dilutes the sample two fold from the original analysis and repeats this step until the internal standards meet the required method criteria. All dilutions on the runlog are listed from the one hundred ml digested. The sample would be originally listed as a 5X. A sample diluted one ml to one liter due to high analyte would be listed as a 1000X. The result would therefore be the dilution factor listed on the runlog times the final volume of one hundred ml times the instrument result divide by the dry weight and the weight. The instrument result in ug/l, final volume units would be in liters, the weight in grams, and the dry weight expressed as a fraction. This yields ug/g which is equivalent to mg/kg.

The waters are digested 100 ml initial volume to 50 ml final volume. The sample is diluted 2.5 fold per the method just prior to analysis. The dilution listed on the runlog is the dilution made from the 50 ml digestate. The result would be the instrument result times the final volume times the dilution factor divided by the initial volume. The result in ug/l is converted to mg/l and reported. All the digestates are left undiluted until analysis to make them as stable as possible. The dilution prior to analysis is specified in the method to reduce damage to the nickel cones.

Method 200.8 does not specify that a CRI, serial dilution, or Interference check be run. Iron, aluminum, calcium, magnesium do not interfere with mass spectroscopy since these masses are at 56 and 57, 27, 40 and 42, and 24 mass units. The only analyte in the method near these masses is manganese at 55 amu. Tungsten is at masses 182 and 184 and only has a small interference from osmium at 182. Both masses were monitored and agreed very well. Had an interference been observed mass 184 would have been used. Either mass may be used for these

samples as no osmium appeared to be present.

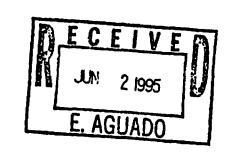
IBA did not perform an idl study for tungsten. A five ug/l ICV was used and recovered very well with low RSD and a SD of 0.016085. From this data the lab reported a PQL of 1 ug/l. Since no linear range study was performed all samples were diluted below the calibration standard of 100 ug/l. CCVs were run at midrange of 50 ug/l.

All samples for this project were stored in the dark since tungsten is light sensitive to insure the stability of the digestates.

Donald Stogner

Inorganics Lab Manager





CLP DATA PACKAGE

SAMPLING DATE APRIL 20, 1995
IEA JOB NO: 20950-51723B-REVISED

VOLUME I OF I

PREPARED BY:

INDUSTRIAL ENVIRONMENTAL ANALYSTS (IEA)

(CERTIFICATION NUMBER 14530)

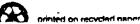
FOR

FOSTER WHEELER ENVIRONMENTAL CORPORATION

PROJECT: CCP

Monroe, Connecticut 203-261-4458 Sunrise, Florida 305:846:1730 Schaumburg. filinois 708-705-0740 N. Billerice, Massachusetts 617:272-5212

Research Triangle Park, North Carolina 919-677-0090



PROJECT: 1254-294

BATCH: 9504608

METHOD: EPA 200.8

Samples:

Twelve (12) Soils and Two (2) Water Samples

The samples were received at Industrial and Environmental Analysts, Inc. (IEA) on April 24, 1995. Each sample was assigned a 9-character "IEA" lab identification number (lab ID) and an abbreviated client ID which is referenced on the IEA Assigned Number Index. All analyses are performed in accordance with EPA approved methodologies and meet the requirements of the IEA Quality Assurance Program. Please see the enclosed data package for your results and Chain of Custody documentation.

The pH of all samples for Metals analysis was less than two (2) at the time of sample preparation.

Any nonconformances associated with the analysis of the samples in this project are as follows:

The quantitation limits for samples 9564668-61 through 65 and 69 through 14 were elevated due to a dilution prior to analysis. The samples were diluted due to high levels of Tungsten.

I certify that this data package is in compliance with the procedures and methods defined for this project, both technically and for completeness, for other than the conditions detailed above. Release of the data contained in this hardcopy data package and in the computer-readable data (if applicable) as submitted has been authorized by the laboratory manager or his designee, as verified by the following signature.

Wallace L. McAnulty

Inorganic Technical Data Reviewer

Wallan I M En

IEA, Inc.

May 80, 1995

IEA, Inc Doct RP768968.NC

000003

CHAIN OF CUSTODY DOCUMENTATION

51723

EBASCO SERVICES INCORPORATED CHAIN OF CUSTODY RECORD

PROJECT	CK.				ERS						/						PRESERVATION
SAMPLERS: (Signature)	DATE	TIME	COMP.	GRAB	NO. CONTAINERS		/3/2 /2/2	5/85/00/00/00/00/00/00/00/00/00/00/00/00/00			/ */ **			/// ¯c	ARIKS IRI' LOCATION	ICED	SPECIFY CHEMICALS ADDED AND FINAL pH IF KNOWN
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EBASCO SERVICES INCORPORATED CHAIN OF CUSTODY RECORD

PROJECT C					VERS					,	/							PRESERVATION
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Print Name	and Company				gnature		· Custody S	cal # (s)	Da	te/Time
Sampled By:										
Received By:							<u> </u>		-	
Relinquished By: John Po.		La	Shill	A	···				4/21/55	1136
Received By: 2 MC COV	med IEA		14U. MY	<u>Cllss</u>					4-24-85	- 1094
Relinquished By:										<u></u>
Received By:	'		<u> </u>							<u>/ c</u>
Mtx = Matrix of Sample. (AI=	Air. AO=Aqueous, LE=Lea	chate	ML=Misc I	iquid MS=	Misc Solids, OI	L, SE=Sedime	ent, SL=Sludge, SC)=Soil)	* Standar	d TAT. 🗗
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IEA OF NEW JERSEY SAMPLE CONTROL CHRONICLE

Sampling Date: 4/1	6175	Job #:	723
Receipt Date: 4/		Job #:	yes
Custody Seal:	Present/Absent Intact/Not Intac	ct Cooler Temp	
Chain of Custody:	Present/Absent		
· · · · · · · · · · · · · · · · · · ·		Preservative	Ck: N
Shipping Bill:	Present Absent	Airbill #:	
Comments:			
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Parameter Sam	ple ID	Parameter	Sample ID
MBAS AMMONIA COD SULFATE NITRATE BOD NITRATE NITRITE RADIUM THORIUM URANIUM Subcontract Lab: Signature:		Prep	1-14
Compositing: :			
Percent Solids:	1-5, [-14	
pH Performed:	-	 .	
Signature:	14	Date: 4-	27-45
Form# SMF00601.N	IJ	Page_ IEA Logbo	OF 98 ok# SM6

TUNGSTEN

:EA Project #: 1254-204

:EA sample #: 9594688-91

:lient Name: IEA - New Jersey

 Matrix: Soil

Date Received: 84/24/95

Date Sampled: 54/28/95

1/ per 12 188.2

arameter	Method	Quantitation Limits	Results	Date Prepared	Date Analyzed Anal	-
Tungsten	EPA 266.8	5.7 mg/kg*	· 51 mg/kg	65/68/95	65/24/95 FW	

mments:

Quantitation limit elevated due to sample dilution prior to analysis. Sample diluted due to high concentration of interferent.

ORM RESP3 Rev. #36994

EA Project #: 1254-264

EA Sample #: 9584688-82

:lient Name: IEA - New Jersey

:lient Proj. I.D.:26958-51723

ample I.D.: CCSS13-51

Matrix: soi

Date Received: \$4/24/95

Date Sampled: \$4/28/95

1/0 paris 3400

'arameter	Method	Quantitation · Limits	Results.	Date Prepared	Date Analyzed	-
-Tungsten	EPA 266.8	59 mg/kg*	1288 mg/kg &10 j	6 5/ 0 8/95	6 5/24/95	

omments:

Quantitation limit elevated due to sample dilution prior to analysis. Sample diluted due to high concentration of interferent.

ORM RESP3 Rev. #36994

80%

Industrial & Environmental Analysts, Inc. (IEA)

Matrix:

MA Project #:

1254-284

SA Sample #:

9564668-83

soil

Lient Wame:

IEA - New Jersey

Date Received: #4/24/95

ilent Proj. I.D.:28958-51723

Date Sampled: \$4/25/95

umple I.D.:

CCS614-81

		Quantitation		Date	Date	
:remeter	Method	Limits	Results	Prepared	Analyzed	Analyst
*************			**********	221		
Tungsten	EPA 200.8	125 mg/kg*	4 3288 mg/kg	T05/08/95	65/24/95	fw

mments:

wantitation limit elevated due to sample dilution prior to analysis. ample diluted due to high concentration of interferent.

RM RESP3 Rev. 836994

IEA Project #: 1254-264

IEA Sample #: 9584658-64

Client Name: IRA - New Jersey

Client Proj. I.D.:28958-51723

Sample I.D.: CCSS15-61.

Matrix: Soil

Date Received: 64/24/95

Date Sampled: \$4/25/95

7.10

Parameter	Method	Quantitation Limits	Results	 Date Analysed	•
r-Tungsten	EPA 288.8		· 3800 mg/kg	65/24/95	

:omments:

Quantitation limit elevated due to sample dilution prior to analysis. Sample diluted due to high concentration of interferent.

ORM RESP3 Rev. #38994

EA Project #: 1254-284

EA Sample #: 9584688-65

Lient Name: IEA - New Jersey

lient Proj. I.D.:28958-51723

soil Matrix:

Date Received: 84/24/95 Date Sampled: \$4/28/95

11

ample I.D.: CC8811-62 .

1/0 ...

arameter	Method	Quantitation Limits	Results	Date Prepared	Date Analyzed Analyst
-Tungsten	EPA 255.8	7.7 mg/kg*		85/88/95	85/24/95 TW

xmments:

mantitation limit elevated due to sample dilution prior to analysis. :ample diluted due to high concentration of interferent.

RM RESP3 Rev. #38994

000014

property of the

CEA Project #: 1254-284

IEA sample #: 9584688-86

Client Name: IEA - New Jersey

:lient Proj. I.D.:28958-51723

sample I.D.:

ccss11-63 /

Matrix: Soil

Date Received: \$4/24/95

Date Sampled: \$4/28/95

Quantitation Date Date

Parameter Method Limits Results Prepared Analyzed Analyst

Chromosome EPA 288.8 8.55 mg/kg 1.4 mg/kg 85/88/95 85/24/95 FW

omments:

'ORM RESP3 Rev. 638994

Matrixs

EA Project #1 1254-254 EA Sample #: 9584688-87

lient Mame:

IEA - New Jersey Date Received: 64/24/95 lient Proj. I.D.:26956-51723 Date Sampled: \$4/28/95

CCFB62-61 ... ample I.D.:

	wakhad	Quantitation	3	Date	Date	
arameter	Method	Limits	Results	Prepared	Analyzed Analyst	
-Tungsten	EPA 266.8	5.861 mg/L	BQL /	65/68/95	65/24/95 FW	•

mments:

)RM RESP3 Rev. #38994

000016 250839

Industrial & Environmental Analysts, Inc. (IEA)

"Matrix:

IEA Project #: 1254-264

IEA Sample #: 9584668-68

Client Name: IEA - New Jersey Client Proj. I.D.:26956-51723 Date Received: 84/24/95 Date Sampled: 64/28/95

Water

Sample I.D.: LTFB0101

		Quantitation		Date	Date	
Parameter	Method	Limits	Results	- .	Analyzed	•
T-Tungsten		6.581 mg/L	BQL 🕜		65/24/95	

Comments:

FORM RESP3 Rev. #30994

EA Project #:

1254-264

EA Sample #:

9584688-89

Matrix:

soil

lient Name:

IEA - New Jersey

Date Received: 84/24/95 Date Sampled: 84/28/95

lient Proj. I.D.:26958-51723 ample I.D.:

LT-8881-81

arameter	Method	Quantitation Limits	Results	Date Prepared	Date Analyzed	•
-Tungsten	EPA 288.8	26# mg/kg*	3888 mg/kg		65/24/95	

Aments:

mantitation limit elevated due to sample dilution prior to analysis. ample diluted due to high concentration of interferent.

RM RESP3 Rev. 636994

EA Project #: 1254-284

EA Sample #: 9584688-15

lient Name: IEA - New Jersey

lient Proj. I.D.:28958-51723

ample I.D.:

LT-8862-61

Matrix:

soil

Date Received: 84/24/95

Date sampled: 84/28/95

756.7

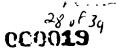
arameter	Method	Quantitation Limits	Results	-	Date Analyzed	=
Tungsten	EPA 266.8				85/24/95	

16200 J

mments:

mantitation limit elevated due to sample dilution prior to analysis. ample diluted due to high concentration of interferent.

RM RESP3 Rev. 636994



4 -

Industrial & Environmental Analysts, Inc. (IEA)

1254-204 SA Project #:

SA Sample 4: 95\$46\$8-11 Matrix:

Lient Name: IEA - New Jersey Date Received: 84/24/95 Date Sampled: 84/28/95

Lient Proj. I.D.:28958-51723 ample I.D.: LT-8883-61

Quantitation Date Date Limits , ... Results Prepared Analyzed Analyst **urameter** Nethod 1288 mg/kg 85/88/95 65/24/95 FW ... Tungsten EPA 200.8 128 mg/kg* 1150 J

mments:

quantitation limit elevated due to sample dilution prior to analysis. sample diluted due to high concentration of interferent.

DRM RESP3 Rev. 838994

Matrix:

A Project #: 1254-264 (Revision)

A sample #: 9554668-12

ient Name: IEA - New Jersey

Date Received: 64/24/95 Date Sampled: 64/26/95

soil

ient Proj. I.D.:26956-51723
mple I.D.: LT-8664-61

rameter	Method	Quantitation Limits	Results	Pate Prepared	Date Analyzed	_
Tungsten	EPA 266.8	258 mg/kg*	4588 ng/kg 4549 J	85/88/95	65/24/95	

mments:

mantitation limit elevated due to sample dilution prior to analysis. ample diluted due to high concentration of interferent.

AM RESP3 Rev. \$38994

Carlo Marin

A Project #: 1254-284

9564668-13 A sample #:

IBA - New Jersey

ient Name: ient Proj. I.D.:28958-51723

LT-8865-61 mple I.D.:

Date Received: 84/24/95

Date Sampled: \$4/25/95

/ ...

rameter	Method	Results	 Date Analysed	-
	EPA 266.8	.1295 mg/kg	85/24/95	

Fores Burney.

11786 - 116

mments:

wantitation limit elevated due to sample dilution prior to analysis. ample diluted due to high concentration of interferent.

RM RESP3 Rev. 638994

EA Project #: 1254-264

EA Sample #: 9564668-14

lient Name: IEA - New Jersey

lient Proj. I.D.:28958-51723

ample I.D.:

LT-6885-81D

Matrix

soil

38 ... T

Date Received: 84/24/95

Date Sampled: \$4/25/95

(13:

Cicco Dand W.

Quantitation Date Analyzed Analyst arameter Method Limits Results Prepared **#5/#8/95** 65/24/95 FW EPA 255.8 146 mg/kg* 1466 mg/kg

omments:

Quantitation limit elevated due to sample dilution prior to analysis. sample diluted due to high concentration of interferent.

ORM RESP3 Rev. #36994

EA Project #: 1254-254

%A Sample #: 9584688 :

lient Name: . . IEA - New Jersey

Matrix: Solid
Date Received: N/A

· lient Proj. I.D.:28958-51723

Date Sampled: N/A

ample I.D.:

QC Blank

i ...

		: : Quantitation	·	Date	Date	_
irameter	Method	Limits	Results	Prepared	Analyzed	Analyst
4492028#228	*************		*********	*******	*******	********
-Tungsten	EPA 200.8	6.58 mg/kg	BQL	65/68/95	65/24/95	T W

uments:

presponding Bamples: 9564688-61 through \$6 and \$9 through 14

RM RESP3 Rev. 838994

Matrix:

IA Project #: 1254-264

A Sample #: 9584688 ... ient Name: IEA - New Jersey

IEA - New Jersey Date Received: N/A

ient Proj. I.D.:28956-51723

Date Sampled: N/A

Water

umple I.D.: QC Blank

1

rameter	Method .		Results	_	:Date	
Tungsten	EPA 266.8	6.561 mg/L	BQL ./	65/68/95	6 5/24/95 I	

mments:

presponding samples: 9584688-97 and 98

RM RESP3 Rev. 030994

Inorganic oc summary Duplicate analysis

IEA Project No.:1254-264
IEA Sample No.: 9564668
Natrix: Soil

IEA Reference No.	Test Parameter	Xethod	DUPLICATE sample (mg/kg)	RESULTS Duplicate (mg/kg)	RPD (%)	Date Analyzed
9584608-12	T-Tungsten	EPA 266.8	_4400 45.4 %	_ 3866 3768	15 ·	85/24/95
		w: न	दम्भू।	- No.	17	

Comments:

Corresponding Samples: 9584688-81 through \$6 and \$9 through 14

FORM IQCSUM Rev 183194

INORGANIC QC SUMMARY DUPLICATE ANALYSIS

IEA Project No.:1254-284
IEA Sample No.: 9584688
Matrix: Water

			DUPLICATE			
 IEA Reference No.	Test Parameter	Hethod	Sample (mg/L)	Duplicate (mg/L)	RPD (%)	Date Analysed
9584688-87	T-Tungsten	EPA 266.8	<6.661 ·	<6.981 ·	6	6 5/24/95

Comments:

Corresponding Samples: 9564688-67 and 68

FORM IQCSUM Rev 183194

INORGANIC QC SUMMARY SPIKE RESULTS

EA Project No.: 1254-254 EA Sample No.: 9554658 Satrix: Soil

:EA	Test		Analysis				
eference No.	Parameter	Method	S A	S R	SSR	♦ R	Date
5\$46\$8-12	T-Tungsten		12	4406 House	3688 2	مندج :	85/24/95

R = (SSR - SR) / (SA) * 100

Control Limits: 75 - 125%

omments:

Percent recovery not calculated due to the sample concentration being greater than four times the concentration of the spiking solution.

orresponding Samples: 9584688-81 through 86 and 89 through 14

· ...

DRM IQCSPK Rev 183194

INORGANIC QC SUMMARY SPIKE RESULTS

IEA Project No.: 1254-264
IEA Sample No.: 9564688
Matrix: Water

			••					
IEA Reference No.		Test Parameter	Nethod	ελ	S R	ssr	€R	Analysis Date
	9584688-87	T-Tungsten		6.10	<6.001	6.16:	185	65/24/95

4R = (6SR - SR)/(SA) * 166

Control Limits: 75 - 125%

Comments:

Corresponding Samples: 9584688-87 and 88

FORM IQCSPK Rev 103194



INORGANIC QC SUMMARY Laboratory Control Sample

IEA Project No.: 1254-284
IEA Sample No.: 9584688

		True Value			Analysis	
Parameter	Method	(mg/L)	Found	* Recovery	Date	
T-Tungsten	EPA 288.8	0.100	5.184	154 '	65/24/95	

make z

Comments:

Control limit is 88% - 128% for all metals.

FORM QCLCSW Rev. 538994

CALIBRATION VERIFICATION

TEA Project No.:

1254-284

CEA Sample No.:

9584668

Parameter	ICV Value (mg/L)	ICV Found (mg/L)	% Recovery	CCV Value (mg/L)	•	t Recovery	CCV End (mg/L)	Recovery	Analysis Date
-Tungsten	6.665	8.665	- "	0.055 0.054	1.1050		سيعهدا.	102	85/24/95

i da√

sals as I letter be

mments:

introl limit is 96% - 116% for all metals, except Hg which is 86% - 126%. introl limit is 85% - 115% for all wet chemistry parameters.

RM CALVER REV #3#994

REFERENCE 13

Pertia.

TO:

M. Heffron

DATE: 8/11/95

FROM:

C. Minch

SUBJECT: CAPTAIN'S COVE SAMPLE RESULTS AND DATA VALIDATION REPORT

Mike,

Enclosed are the results and data validation report for the uranium and thorium analyses conducted on samples collected from the Captain's Cove Site. Edgar asked me to send them directly to you because of the time constraints. If you have any questions, I can be reached at (908) 270 - 0988.

Celia

cc: E. Aguado



UNITED STATES ENVIRONMENTAL PROTECTION AGENCY OFFICE OF RADIATION AND INDOOR AIR

National Air and Radiation Environmental Laboratory 540 South Morris Avenue, Montgomery, AL 36115-2601 (334) 270-3400

August 4, 1995

MEMORANDUM

SUBJECT: Radiochemical Results for

Captain's Cove Samples

FROM:

John Griggs, Chief John / Lu

Monitoring and Analytical Services Branch

Vicki Lloyd, Director

NAREL Technical Support Center (TSC)

TO:

Catherine Moyik, Site Assessment Manager

Superfund, Region 2

Attached are data packages for gross alpha and beta, gamma and isotopic uranium and thorium analyses of water and soil samples collected at the Captain's Cove Condominium Site located in Glen Cove, Nassau County, New York. The samples constitute NAREL batch numbers 95-00015 and 95-00016.

Although no analytical problems were encountered in analyzing NAREL Sample T34C 95.03174, we are reanalyzing the sample because of possible inconsistencies between the measured activities of radionuclides which are normally in equilibrium. The results of the reanalysis will be reported as soon as they are available.

Radiochemical analyses usually require the subtraction of an instrument background measurement from a gross sample measurement. Both values are positive, but when the sample activity is low, random variations in the two measurements can cause the gross value to be less than the background, resulting in a measured activity less than zero. Although negative activities have no physical significance, they do have statistical significance, as for example in the evaluation of trends or the comparison of two groups of samples.

For all analyses except gamma spectroscopy, it is the policy of NAREL to report results as generated, whether positive, negative, or zero, together with the 2-sigma measurement uncertainty and a sample-specific estimate of the minimum detectable concentration (MDC). The activity, uncertainty, and MDC are given in the same units. The activity and 2-sigma uncertainty for a radionuclide measured by gamma spectroscopy are

reported only if the nuclide is detected; so, the results of gamma analyses are never zero or negative. Nuclides that are not detected do not appear in the report, with the exception of Ba-140, Cs-137, I-131, K-40, Ra-226, and Ra-228. If one of these six nuclides is undetected, NAREL reports it as "Not Detected," or "ND," and provides a sample-specific estimate of the MDC.

Specific information concerning all aspects of the radiological analysis of the samples is contained in the batch case narratives of the data packages. If you have any questions concerning the analytical results, the analytical process, or the reporting format, contact Dr. John Griggs at (334) 270-3450. If you have any project-specific questions or questions concerning data application, contact Vicki Lloyd at (334) 270-3467.

Attachments

cc: Paul Giardina, Region 2, w/o attachments
Edgar Aguado, Ebasco, w/attachments
Mary Clark, (6601J), w/o attachments
Sam T. Windham

CAPTAIN'S COVE
RADIOCHEMICAL
DATA VALIDATION REPORT
NAREL BATCH # 95-00015

Prepared by:	Date:	
Coule 7? March	8/11/95	_
Cecelia N. Minch		_

CAPTAINS COVE DATA VALIDATION REPORT NAREL Batch # 95-00015

SUMMARY:

This case consisted of 12 soil samples collected on April 20, 1995 and designated for the analysis of uranium and thorium by alpha spectrometry. One field duplicate pair (LTSS05-01/05D) was collected and analyzed with satisfactory results.

The laboratory documented in the narrative that problems were encountered with the uranium analyses of samples LT-SS03-01.

LT-SS04-01 and the replicate of LT-SS04-01. Matrix interferences may be responsible since reanalyses produced similar results. The data user should also be aware that the laboratory thought it prudent to reanalyze sample CC-SS13-01 due to possible inconsistencies (see letter of August 4, 1995), the results of which will follow at a later date.

All data were evaluated for Level D DQO. employing USEPA Region II inorganic data validation criteria to the extent possible. The specifics for each parameter and associated QC are detailed below. The sample identifications used in this report have been truncated for expediency. Unless otherwise indicated, all sample IDs are suffixed with -01.

All data are considered acceptable and valid with the following qualifications.

HOLDING TIMES:

A holding time of 180 days has been applied to the samples. All samples were analyzed within this holding time.

TRACERS:

The following samples exhibited low tracer recovery (<80%). As a result, all reported values for the associated isotopes may be biased low and, therefore, were qualified as estimated (J).

Uranium: SS03; SS04

thorium: SS05, SS05D, SS13, SS14, SS15

The replicate of SS04/also experienced low recovery, but no action was necessary.

MATRIX SPIKES:

The soil matrix spike duplicate exceeded recovery criteria and the RPD for U-235. Consequently, U-235 sample results greater than the MDC may be biased high and were qualified as estimated (J) as follows:

Qualified "J":

SS05, SS05D, SS12, SS14, SS11-02, SS11-03

Samples SS01. SS03 and SS04 would also have been qualified "J" for U-235, but were previously qualified for other criteria.

The recovery of Th-230 was acceptable.

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REPLICATES:

Two replicate pairs were analyzed and evaluated for reproducibility. The data were qualified as estimated (J) only when the RPD exceeded 50% and the results reported for both analyses were greater than the MDC. The direction of bias in this instance is unknown.

Qualified "J":

Th-228: SS01, SS02, SS03, SS04, SS11-02, SS11-03 Th-232: SS01, SS02, SS03, SS04, SS11-02, SS11-03

Both sets of replicate data were acceptable for uranium.

FIELD DUPLICATE:

The results of the field duplicate were acceptable.

BLANK CONTAMINATION:

No qualifications were required.

INSTRUMENT CALIBRATION:

An efficiency check standard was analyzed on each detector approximately every 7 days, the results of which were plotted on a control chart. The values obtained were evaluated for compliance with the ± 2 standard deviation limits defined on the charts. Sample analyses bracketed by acceptable standards are deemed acceptable. However, the standards which followed 2 samples were not within the established limits. In both cases, any sample values greater than the MDC were qualified as estimated (J) and may be biased low. The following data were qualified "J":

SS01: U-234, U-235, U238 V SS12: Th-228, Th-230, Th-232

(2et. 13) 7 of 45

CASE NARRATIVE

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

NAREL Batch #:

95-00015

I. RECEIPT

A. Sample Information

NAREL Sample ID	Client Sample ID	Sample Matrix	Date Collected	Date Received	Date Analyzed
<u></u>					
T34C 95.03167	LT-SS01-01	Soil	04/20/95	04/21/95	07/07/95
T34C 95.03168	LT-SS02-01	Soil	04/20/95	04/21/95	06/28/95
£34C 95.03169	£T-SS03-01	Soil	04/20/95	04/21/95	06/26/95
T34C 95.03170	ALT-SS04-01	Soil	04/20/95	04/21/95	06/26/95
T34C 95.03171	LT-SS05-01	Soil	04/20/95	04/21/95	06/28/95
T34C 95.03172	LT-SS05-01D	Soil	04/20/95	04/21/95	06/28/95
T34C 95.03173	CC-SS12-01	Soil	04/20/95	04/21/95	07/07/95
T34C 95.03174	CC-SS13-01	Soil	04/20/95	04/21/95	06/28/95
T34C 95.03175	CC-SS14-01	Soil	04/20/95	04/21/95	06/28/95
T34C 95.03176	CC-SS15-01	Soil	04/20/95	04/21/95	06/28/95
T34C 95.03177	CC-SS11-02	Soil	04/20/95	04/21/95	06/26/95
T34C 95.03178	CC-SS11-03	Soil	04/20/95	04/21/95	06/26/95

B. Documentation

Exceptions:

No exceptions were encountered.

II. ANALYSIS

A. Holding Times:

All holding times were met.

B. Preparation

Exceptions:

No exceptions were encountered.

C. Analytical

Exceptions:

cNAREL samples T34C 95.3169, T34C 95.3170 and T34C 95.3170X formed a purple precipitate during the coprecipitation step of the uranium procedure. This step normally results in an essentially "massless" sample being deposited onto a planchet. Because of the presence of the precipitates in these samples, the alpha spectra contain smeared peaks which were rejected by NAREL counting room data reviewers. The samples were reanalyzed and

cimilar results were obtained. We believe these camples contain interferences which cause the formation of a problematic amount of precipitate during the coprecipitation step. The results of the priginal analyses are contained in this report. We recommend that the results be used only as a finalitative means of indicating the presence of these aradionuclides and not as a quantitative measure of their concentration and that the results of the replicate analysis of sample T34C 95.3170 not be used in the evaluation of the quality control samples.

The uranium analyses on NAREL samples T34C 95.03167 and T34C 95.03173 gave measured yields greater than 104%. These two samples were recounted and the results from the recounts are provided in this package.

The result from the efficiency check for detector AS10 on 6/26/95 was lost, and the check was not repeated. No samples in this batch were analyzed on AS10.

Detector AS28 is not currently in operation.

III. QUALITY CONTROL

A. Reagent Blank:

All associated reagent blanks met NAREL QC

criteria.

B. Tracer Yields:

All samples met NAREL QC limits.

C. Matrix Spike:

All spike recoveries were within NAREL QC

limits.

D. Replicate Results:

Although the results of the replicate analysis of sample T34C 95.3170 are provided in this report, we recommend that the results of the replicate analysis of this sample not be used in the evaluation of the quality control samples. The analytical problems associated with this sample are described in the Analytical Exceptions section of the case narrative.

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CASE NARRATIVE

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

NAREL Batch #:

95-00015

I. RECEIPT

Sample Information A.

NAREL	Client	Sample	Date	Date Received	Date
Sample ID	Sample ID	Matrix	Collected		Analyzed
T34C 95.03167 T34C 95.03168 T34C 95.03169 T34C 95.03170 T34C 95.03171 T34C 95.03172 T34C 95.03174 T34C 95.03175 T34C 95.03176 T34C 95.03177 T34C 95.03177	LT-SS01-01 LT-SS02-01 LT-SS03-01 LT-SS05-01 LT-SS05-01D CC-SS12-01 CC-SS13-01 CC-SS15-01 CC-SS15-01 CC-SS11-02 CC-SS11-03	Soil	04/20/95 04/20/95 04/20/95 04/20/95 04/20/95 04/20/95 04/20/95 04/20/95 04/20/95 04/20/95	04/21/95 04/21/95 04/21/95 04/21/95 04/21/95 04/21/95 04/21/95 04/21/95 04/21/95 04/21/95 04/21/95	07/07/95 06/28/95 06/26/95 06/26/95 06/28/95 06/28/95 07/07/95 06/28/95 06/28/95 06/28/95 06/26/95

B. Documentation Exceptions:

No exceptions were encountered.

II. **ANALYSIS**

A. Holding Times:

All holding times were met.

B. Preparation

Exceptions:

No exceptions were encountered.

C. Analytical Exceptions:

eNAREL samples T34C 95.3169, T34C 95.3170 and T34C 95.3170X formed a purple precipitate during

the coprecipitation step of the uranium procedure.

This step normally results in an essentially

"massless" sample being deposited onto a planchet. Because of the presence of the precipitates in these samples, the alpha spectra contain smeared peaks which were rejected by NAREL counting room data

reviewers. The samples were reanalyzed and

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URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: Reagent Blank

NAREL Sample #:

RBLK 95.03179

NAREL Batch #: 95-00015

Date Collected:

<u>05/03/95</u>

Matrix:

<u>Soil</u>

Date Received:

05/03/95

Wet weight:

<u>N/A</u>

Date Analyzed:

06/26/95

Dry weight:

N/A

Analyst:

<u> AS</u>

Ash weight:

N/A

Method:

EERF-00.06

Vol/Wt Prepared:

N/A

Detector ID:

<u>AS25</u>

Activity units:

pCi/Samp

Analytical Results

Nuclide	Activity	20 Uncertainty	MDC
U-234 -	4.27E-02	± 2.47E-02	3.12E-02
U-235	1.02E-03	± 6.62E-03	2.30E-02
U-238	3.45E-02	± 2.62E-02	4.46E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

Comments:

12

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS01-01

NAREL Sample #:

T34C 95.03167

NAREL Batch #: 95-00015

Date Collected:

<u>04/20/95</u>

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

530.0 g

Date Analyzed:

<u>07/07/95</u>

Dry weight:

402.6 g

Analyst:

<u> AS</u>

Ash weight:

358.3 g

Method:

EERF-00.06

Vol/Wt Prepared:

0.0051 gash

Detector ID:

<u>AS17</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	4.63E+01 ブ	± 8.72E+00	3.19E+00
U-235	2.11E+00	± 1.70E+00	1.35E+00
U-238	5.22E+01	± 9.34E+00	3.41E+00

In June 1 Barriage Tour **QA/QC Reference Samples**

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

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URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: <u>LT-SS02-01</u>

NAREL Sample #:

T34C 95.03168

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

539.5 g

Date Analyzed:

06/28/95

Dry weight:

467.3 g

Analyst:

<u> AS</u>

Ash weight:

421.5 g

Method:

EERF-00.06

Vol/Wt Prepared:

0.0050 gash

Detector ID:

AS12

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	1.07E+01	± 4.94E+00	5.75E+00
U-235	9.47E-01	± 1.32E+00	1.82E+00
U-238	2.62E+01	± 7.44E+00	4.87E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name: Captain's Cove Client Sample ID: <u>LT-SS03-01</u>

NAREL Sample #:

T34C 95.03169

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

932.5 g

Date Analyzed:

06/26/95

Dry weight:

768.1 g

Analyst:

<u>AS</u>

Ash weight:

703.6 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2541 gash

Detector ID:

AS07

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	26 Uncertainty	MDC
U-234	3.28E+00 J	± 4.85E-01	8.25E-02
U-235	7.19E-01	± 1.89E-01	7.07E-02
U-238	3.24E+00 ✓	± 4.83E-01	8.25E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

14 of 45

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS04-01

NAREL Sample #:

T34C 95.03170

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

586.9 g

Date Analyzed:

06/26/95

Dry weight:

459.2 g

Analyst:

<u>AS</u>

Ash weight:

416.9 g

Method:

EERF-00.06

Vol/Wt Prepared:

0.2536 gash

Detector ID:

AS09

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	1.76E+00 J	± 3.89E-01	1.34E-01
U-235	5.24E-01	± 1.91E-01	1.03E-01
U-238	9.62E-01	± 2.70E-01	1.21E-01

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS04-01

NAREL Sample #:

T34C 95.03170X

NAREL Batch #: 95-00015

Date Collected:

<u>04/20/95</u>

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

586.9 g

Date Analyzed:

06/26/95

Dry weight:

459.2 g

Analyst:

<u>AS</u>

Ash weight:

416.9 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2507 gash

Detector ID:

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	1.79E+00	± 4.01E-01	1.51E-01
U-235	6.44E-01	± 2.17E-01	1.07E-01
U-238	9.90E-01	± 2.79E-01	1.25E-01

QA/QC Reference Samples

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

Comments:

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URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS05-01

NAREL Sample #:

T34C 95.03171

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

406.8 g

Date Analyzed:

06/28/95

Dry weight:

293.6 g

Analyst:

<u>AS</u>

Ash weight:

242.5 g

Method:

EERF-00.06

Vol/Wt Prepared:

0.0050 gash

Detector ID:

AS21

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	26 Uncertainty	MDC
U-234	1.55E+02	± 1.78E+01	2.97E+00 .
U-235	5.31E+00 J	± 2.59E+00	2.30E+00
U-238	1.65E+02	± 1.86E+01	3.30E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

Comments:

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URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS05-01D

NAREL Sample #:

T34C 95.03172

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

408.3 g

Date Analyzed:

06/28/95

Dry weight:

294.6 g

Analyst:

<u>AS</u>

Ash weight:

242.2 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0051 gash

Detector ID:

AS22

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	1.48E+02	± 1.78E+01	2.70E+00
U-235	5.82E+00 エ	± 2.72E+00	1.24E+00
U-238	1.54E+02	± 1.83E+01	2.70E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
. Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

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URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS12-01

NAREL Sample #:

T34C 95.03173

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

444.6 g

Date Analyzed:

07/07/95

Dry weight:

402.2 g

Analyst:

<u> AS</u>

Ash weight:

386.9 g

Method:

EERF-00.06

Vol/Wt Prepared:

0.2517 gash

Detector ID:

<u>AS18</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	20 Uncertainty	MDC
U-234	7.77E-01	± 1.55E-01	4.45E-02
U-235	3.45E-02 丁	± 3.03E-02	2.66E-02
U-238	7.11E-01	± 1.47E-01	2.66E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS12-01

NAREL Sample #:

T34C 95.03173M

NAREL Batch #: 95-00015

Date Collected:

<u>04/20/95</u>

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

444.6 g

Date Analyzed:

06/26/95

Dry weight:

402.2 g

Analyst:

<u> AS</u>

Ash weight:

386.9 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2502 gash

Detector ID:

AS17

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	4.14E+00	± 4.72E-01	6.37E-02
U-235	2.23E-01	± 8.55E-02	7.37E-02
U-238	3.98E+00	± 4.59E-01	7.79E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

20 of 45

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS12-01

NAREL Sample #:

T34C 95.03173S

NAREL Batch #: 95-00015

Date Collected:

<u>04/20/95</u>

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

444.6 g

Date Analyzed:

<u>06/26/95</u>

Dry weight:

402.2 g

Analyst:

<u>AS</u>

Ash weight:

386.9 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2517 gash

Detector ID:

AS18

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	4.23E+00	± 4.44E-01	4.19E-02
U-235	2.86E-01	± 8.70E-02	2.50E-02
U-238	4.00E+00	± 4.27E-01	4.19E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

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URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS13-01

NAREL Sample #:

T34C 95.03174

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

490.8 g

Date Analyzed:

06/28/95

Dry weight:

418.0 g

Analyst:

<u> AS</u>

Ash weight:

398.8 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0252 gash

Detector ID:

AS23

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	1.11E+00	± 6.15E-01	7.58E-01
U-235 <	2.47E-02	± 1.60E-01	5.58E-01
U-238	3.45E-01	± 3.70E-01	6.72E-01

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

22 of 45

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS14-01

NAREL Sample #:

T34C 95.03175

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

453.4 g

Date Analyzed:

06/28/95

Dry weight:

366.5 g

Analyst:

<u> AS</u>

Ash weight:

351.1 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0126 gash

Detector ID:

AS24

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	2.39E+01	± 3.91E+00	1.01E+00
U-235	1.07E+00 丁	± 7.43E-01	5.13E-01
U-238	1.86E+01	± 3.37E+00	5.13E-01

QA/QC Reference Samples

QC Sample	NAREL Sample Number
Reagent Blank	RBLK 95.03179
Replicate 1	T34C 95.03170X
Replicate 2	T34C 95.03177X
Matrix Spike	T34C 95.03173M
Matrix Spike Duplicate	T34C 95.03173S

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URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: <u>CC-SS15-01</u>

NAREL Sample #:

T34C 95.03176

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

459.7 g

Date Analyzed:

06/28/95

Dry weight:

376.3 g

Analyst:

<u> AS</u>

Ash weight:

359.0 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0126 gash

Detector ID:

AS25

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	2.08E+01	± 4.64E+00	1.83E+00
U-235	5.10E-01	± 7.71E-01	1.65E+00
U-238	1.84E+01	± 4.46E+00	3.27E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

24 of 45

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS11-02

NAREL Sample #:

T34C 95.03177

NAREL Batch #: 95-00015

Date Collected:

<u>04/20/95</u>

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

272.9 g

Date Analyzed:

06/26/95

Dry weight:

186.2 g

Analyst:

<u> AS</u>

Ash weight:

131.4 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2533 gash

Detector ID:

<u>AS19</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	26 Uncertainty	MDC
U-234	8.47E-01	± 1.55E-01	5.28E-02
U-235	3.88E-02 ブ	± 3.06E-02	3.76E-02
U-238	9.53E-01	± 1.66E-01	5.28E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS11-02

NAREL Sample #:

T34C 95.03177X

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

272.9 g

Date Analyzed:

06/26/95

Dry weight:

186.2 g

Analyst:

<u>AS</u>

Ash weight:

131.4 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2517 gash

Detector ID:

AS20

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
U-234	7.48E-01	± 1.47E-01	5.24E-02
U-235	4.18E-02	± 3.30E-02	4.04E-02
U-238	7.13E-01	± 1.44E-01	4.72E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

Comments:

FORM-1 URANTUM

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URANIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS11-03

NAREL Sample #:

T34C 95.03178

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

361.6 g

Date Analyzed:

06/26/95

Dry weight:

318.4 g

Analyst:

<u> AS</u>

Ash weight:

303.1 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2526 gash

Detector ID:

<u>AS24</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	20 Uncertainty	MDC
U-234	5.05E-01	± 1.22E-01	5.19E-02
U-235	2.75E-02 =	± 2.70E-02	2.65E-02
U-238	4.54E-01	± 1.14E-01	2.65E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

CASE NARRATIVE

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

NAREL Batch #:

95-00015

I. RECEIPT

A. Sample Information

NAREL	Client	Sample	Date	Date	Date
Sample ID	Sample ID	Matrix	Collected	Received	Analyzed
T34C 95.03167	LT-SS01-01	Soil	04/20/95	04/21/95	06/29/95
T34C 95.03168	LT-SS02-01	Soil	04/20/95	04/21/95	06/29/95
T34C 95.03169	LT-SS03-01 .	Soil	04/20/95	04/21/95	06/27/95
T34C 95.03170	LT-SS04-01	Soil	04/20/95	04/21/95	06/27/95
T34C 95.03171	LT-SS05-01	Soil	04/20/95	04/21/95	06/27/95
T34C 95.03172	LT-SS05-01D	Soil	04/20/95	04/21/95	06/27/95
T34C 95.03173	CC-SS12-01	Soil	04/20/95	04/21/95	06/27/95
T34C 95.03174	CC-SS13-01	Soil	04/20/95	04/21/95	06/27/95
T34C 95.03175	CC-SS14-01	Soil	04/20/95	04/21/95	06/28/95
T34C 95.03176	CC-SS15-01	Soil	04/20/95	04/21/95	06/28/95
T34C 95.03177	CC-SS11-02	Soil	04/20/95	04/21/95	06/27/95
T34C 95.03178	CC-SS11-03	Soil	04/20/95	04/21/95	06/27/95

B. Documentation

Exceptions:

No exceptions were encountered.

II. ANALYSIS

A. Holding Times:

All holding times were met.

B. Preparation

Exceptions:

No exceptions were encountered.

C. Analytical

Exceptions:

The results from the efficiency check for detector AS10 on 6/26/95 was lost, and the check was not repeated. No samples in this batch were analyzed

on detector AS10.

Detector AS28 is not currently in operation.

met.15

THORIUM 28 of 45 NAREL Batch # 95-00015 PAGE 2

III. QUALITY CONTROL

A. Reagent Blank:

All associated reagent blanks met NAREL QC

criteria.

B. Tracer Yields:

All samples met NAREL QC limits.

C. Matrix Spike:

All spike recoveries were within NAREL QC

limits.

D. Replicate Results:

The results of the replicate analysis on NAREL

cample 95.03177 did not meet NAREL's

acceptance criteria. «

IV. I certify that this data package complies with the terms and conditions of the Quality Assurance Project Plan, both technically and for completeness, other than the exceptions detailed above. Release of the data contained in this package has been authorized by the Chief of the Monitoring and Analytical Services Branch and the NAREL Quality Assurance Coordinator, or their designees, as verified by the following signatures.

ames B. Moore

Quality Assurance Coordinator

Date

8/4/95

Data

Chief, Monitoring

and Analytical Services Branch

29 0445

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: Reagent Blank

NAREL Sample #:

RBLK 95.03179

NAREL Batch #: 95-00015

Date Collected:

05/03/95

Matrix:

<u>Soil</u>

Date Received:

<u>05/03/95</u>

Wet weight:

N/A

Date Analyzed:

<u>06/27/95</u>

Dry weight:

N/A

Analyst:

<u>AS</u>

Ash weight:

N/A

Method:

EERF-00.06

Vol/Wt Prepared:

N/A

Detector ID:

<u>AS21</u>

Activity units:

pCi/Samp

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	6.47E-03	± 1.92E-02	3.85E-02
Th-228	-2.22E-02	± 2.55E-02	5.22E-02
Th-230	1.02E-02	± 9.81E-03	1.33E-02
Th-232 <	5.39E-03	± 8.56E-03	1.50E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: <u>LT-SS01-01</u>

NAREL Sample #:

T34C 95.03167

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

530.0 g

Date Analyzed:

06/29/95

Dry weight:

402.6 g

Analyst:

<u>AS</u>

Ash weight:

358.3 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0051 gash

Detector ID:

AS29

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	5.19E+00	± 3.87E+00	4.42E+00
Th-228	4.49E+00 J	± 5.24E+00	8.62E+00
Th-230	1.11E+01	± 3.70E+00	1.57E+00
Th-232	1.01E+01 J	± 3.55E+00	1.85E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS02-01

NAREL Sample #:

T34C 95.03168

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

539.5 g

Date Analyzed:

06/29/95

Dry weight:

467.3 g

Analyst:

<u> AS</u>

Ash weight:

421.5 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0050 gash

Detector ID:

AS30

Activity units:

pCi/gdrv

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	-1.97E-01	± 2.79E+00	6.52E+00
Th-228	1.27E+01 ゴ	± 6.27E+00	8.94E+00
Th-230	2.00E+01	± 4.96E+00	1.84E+00
Th-232	1.77E+01 J	± 4.65E+00	1.57E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS03-01

NAREL Sample #:

T34C 95.03169

NAREL Batch #: 95-00015

Date Collected:

<u>04/20/95</u>

Matrix:

Soil

Date Received:

<u>04/21/95</u>

Wet weight:

932.5 g

Date Analyzed:

<u>06/27/95</u>

Dry weight:

768.1 g

Analyst:

<u> AS</u>

Ash weight:

703.6 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2541 gash

Detector ID:

AS06

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	4.84E-01	± 1.83E-01	1.21E-01
Th-228	3.12E+00 J	± 3.32E-01	1.98E-01
Th-230	5.38E+00	± 4.19E-01	9.69E-02
Th-232	3.21E+00 5	± 3.28E-01	1.56E-01

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS04-01

NAREL Sample #:

T34C 95.03170

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

586.9 g

Date Analyzed:

<u>06/27/95</u>

Dry weight:

459.2 g

Analyst:

<u>AS</u>

Ash weight:

416.9 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2536 gash

Detector ID:

AS07

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	1.04E-01	± 1.09E-01	1.66E-01
Th-228	6.66E-01 J	± 1.84E-01	2.06E-01
Th-230	9.25E-01	± 1.74E-01	4.90E-02
Th-232	5.83E-01 J	± 1.38E-01	4.17E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS04-01

NAREL Sample #:

T34C 95.03170X

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

586.9 g

Date Analyzed:

06/27/95

Dry weight:

459.2 g

Analyst:

<u>AS</u>

Ash weight:

416.9 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2507 gash

Detector ID:

AS09

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	20 Uncertainty	MDC
Th-227	1.71E-01	± 1.25E-01	1.25E-01
Th-228	7.18E-01	± 2.08E-01	2.32E-01
Th-230	1.22E+00	± 2.18E-01	6.48E-02
Th-232	5.71E-01	± 1.50E-01	6.48E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
- Matrix Spike Duplicate	T34C 95.03173S	

Comments:

QC ARE

NOT

QUALIFIED

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS05-01

NAREL Sample #:

T34C 95.03171

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

406.8 g

Date Analyzed:

<u>06/27/95</u>

Dry weight:

293.6 g

Analyst:

<u>AS</u>

Ash weight:

242.5 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0050 gash

Detector ID:

<u>AS27</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	2.18E+01 J	± 9.46E+00	8.83E+00
Th-228	3.45E+01	± 9.08E+00	9.59E+00
Th-230	3.44E+02	± 2.47E+01	2.45E+00
Th-232	2.48E+01	± 6.49E+00	3.54E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

Comments:

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FORM-1 THORIUM

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: LT-SS05-01D

NAREL Sample #:

T34C 95.03172

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

<u>04/21/95</u>

Wet weight:

408.3 g

Date Analyzed:

06/27/95

Dry weight:

294.6 g

Analyst:

<u>AS</u>

Ash weight:

242.2 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0051 gash

Detector ID:

<u>AS29</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	20 Uncertainty	MDC
Th-227	2.75E+01 J	± 1.15E+01	8.22E+00
Th-228	3.34E+01	± 1.00E+01	1.05E+01
Th-230	3.03E+02	± 2.60E+01	2.70E+00
Th-232	2.51E+01	± 7.32E+00	3.17E+00 ·

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS12-01

NAREL Sample #:

T34C 95.03173

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

444.6 g

Date Analyzed:

06/27/95

Dry weight:

402.2 g

Analyst:

<u>AS</u>

Ash weight:

<u>386.9 g</u>

Method:

EERF-00.06

Vol/Wt Prepared: 0.2517 gash

Detector ID:

<u>AS11</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	3.47E-02	± 1.03E-01	2.06E-01
Th-228	6.78E-01 J	± 2.02E-01	2.27E-01
Th-230	6.13E-01 J	± 1.56E-01	7.11E-02
Th-232	8.93E-01 =	± 1.88E-01	7.11E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY 38 6245

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS12-01

NAREL Sample #:

T34C 95.03173M

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

<u>04/21/95</u>

Wet weight:

444.6 g

Date Analyzed:

<u>06/27/95</u>

Dry weight:

402.2 g

Analyst:

<u> AS</u>

Ash weight:

386.9 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2502 gash

Detector ID:

AS12

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	0.00E+00	± 7.23E-02	1.85E-01
Th-228	1.10E+00	± 2.48E-01	2.37E-01
Th-230	2.48E+01	± 1.12E+00	9.04E-02
Th-232	1.24E+00	± 2.29E-01	7.58E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

Comments:

QC AKE

NOT

QUALIFIED

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY 39 of 45

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS12-01

NAREL Sample #:

T34C 95.03173S

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

444.6 g

Date Analyzed:

06/27/95

Dry weight:

402.2 g

Analyst:

<u>AS</u>

Ash weight:

386.9 g

Method:

EERF-00.06

Vol/Wt Prepared:

0.2517 gash

Detector ID:

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	1.14E-01	± 1.14E-01	1.70E-01
Th-228	1.56E+00	± 2.53E-01	2.14E-01
Th-230	2.44E+01	± 9.98E-01	5.85E-02
Th-232	1.63E+00	± 2.32E-01	8.39E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

Comments:

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UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY 40 0 45

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS13-01

NAREL Sample #:

T34C 95.03174

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

490.8 g

Date Analyzed:

<u>06/27/95</u>

Dry weight:

418.0 g

Analyst:

<u> AS</u>

Ash weight:

398.8 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0252 gash

Detector ID:

<u>AS30</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	1.43E+00 J	± 1.71E+00	2.76E+00 ·
Th-228	5.64E-02	± 1.33E+00	2.53E+00
Th-230	6.30E-01	± 5.90E-01	7.21E-01
Th-232	7.92E-02	± 2.49E-01	6.14E-01

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY 41 of 45

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS14-01

NAREL Sample #:

T34C 95.03175

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

453.4 g

Date Analyzed:

06/28/95

Dry weight:

366.5 g

Analyst:

<u>AS</u>

Ash weight:

351.1 g

Method:

EERF-00.06

Vol/Wt Prepared:

0.0126 gash

Detector ID:

AS06

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	20 Uncertainty	MDC
Th-227	4.87E+00 丁	± 3.90E+00	4.69E+00
Th-228	1.94E+01	± 5.66E+00	5.91E+00
Th-230	4.52E+01	± 7.47E+00	3.68E+00
Th-232	2.00E+01 ↓	± 5.71E+00	5.90E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

Comments:

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UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY 42 & 45

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS15-01

NAREL Sample #:

T34C 95.03176

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

459.7 g

Date Analyzed:

06/28/95

Dry weight:

376.3 g

Analyst:

<u>AS</u>

Ash weight:

359.0 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.0126 gash

Detector ID:

AS07

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	20 Uncertainty	MDC
Th-227	2.61E+00 J	± 3.69E+00	6.25E+00
Th-228	1.95E+01	± 5.79E+00	6.13E+00
Th-230	3.94E+01	± 6.90E+00	1.80E+00
Th-232	1.62E+01 ↓	± 4.40E+00	1.53E+00

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY 43 545

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS11-02

NAREL Sample #:

T34C 95.03177.

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

Soil

Date Received:

04/21/95

Wet weight:

272.9 g

Date Analyzed:

<u>06/27/95</u>

Dry weight:

186.2 g

Analyst:

<u>AS</u>

Ash weight:

131.4 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2533 gash

Detector ID:

<u>AS18</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	2.36E-01	± 1.05E-01	9.30E-02
Th-228	1.13E+00 J	± 1.65E-01	1.34E-01
Th-230	9.26E-01	± 1.33E-01	2.43E-02
Th-232	1.22E+00 J	± 1.53E-01	1.41E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

Ref 13 UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY 44 64 45

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS11-02

NAREL Sample #:

T34C 95.03177X

NAREL Batch #: 95-00015

Date Collected:

04/20/95

Matrix:

<u>Soil</u>

Date Received:

<u>04/21/95</u>

Wet weight:

272.9 g

Date Analyzed:

06/27/95

Dry weight:

186.2 g

Analyst:

<u>AS</u>

Ash weight:

131.4 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2517 gash

Detector ID:

AS19

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	MDC
Th-227	1.07E-01	± 9.24E-02	1.28E-01
Th-228	3.81E-01	± 1.30E-01	1.60E-01
Th-230	7.70E-01	± 1.40E-01	4.57E-02
Th-232	5.36E-01	± 1.17E-01	4.57E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

Comments:

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UNITED STATES ENVIRONMENTAL PROTECTION AGENCY NATIONAL AIR AND RADIATION ENVIRONMENTAL LABORATORY

THORIUM RADIOCHEMICAL ANALYTICAL RESULTS

Project Name:

Captain's Cove

Client Sample ID: CC-SS11-03

NAREL Sample #:

T34C 95.03178

NAREL Batch #: 95-00015

Date Collected:

<u>04/20/95</u>

Matrix:

<u>Soil</u>

Date Received:

04/21/95

Wet weight:

361.6 g

Date Analyzed:

06/27/95

Dry weight:

318.4 g

Analyst:

<u>AS</u>

Ash weight:

303.1 g

Method:

EERF-00.06

Vol/Wt Prepared: 0.2526 gash

Detector ID:

<u>AS20</u>

Activity units:

pCi/gdry

Analytical Results

Nuclide	Activity	2σ Uncertainty	· MDC
Th-227	-3.05E-02	± 5.57E-02	1.62E-01
Th-228	4.71E-01 5	± 1.62E-01	2.08E-01
Th-230	4.67E-01	± 1.16E-01	4.81E-02
Th-232	4.45E-01 J	± 1.14E-01	4.81E-02

QA/QC Reference Samples

QC Sample	NAREL Sample Number	
Reagent Blank	RBLK 95.03179	
Replicate 1	T34C 95.03170X	
Replicate 2	T34C 95.03177X	
Matrix Spike	T34C 95.03173M	
Matrix Spike Duplicate	T34C 95.03173S	

REFERENCE 14

AND APPLIED PHYSICS ULTING EDITOR

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THE ATOMIC NUCLEUS

Robley D. Evans, Ph.D.

PROFESSOR OF PHYSICS
MASSACHUSETTS INSTITUTE OF TECHNOLOGY

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r of the series from its inception in as Consulting Editor from 1939 to

New York Toronto London

McGRAW-HILL BOOK COMPANY, INC.

1955

(b) What is the "partial half-period" for negatron β decay, i.e., the halfperiod for a Cu44 nucleus in which the possibility of other modes of decay has been

(c) Evaluate in millicuries a source of Cu⁴⁴ which emits 3.7 × 10⁷ negatron

β rays per second.

Ans.: (b) 32.0 hr; (c) 2.5 mc. 2. Determine the half-period of K⁴⁰, knowing that ordinary potassium (a) is a mixture of K39, K40, and K41 containing 0.0119 atom per cent K40; (b) emits 31 β rays/sec per gram in transitions of $K^{40} \xrightarrow{\beta} Ca^{40}$; and (c) emits 3.4 γ rays/sec per gram in electron-capture transitions $K^{40} \xrightarrow{BC} A^{40}$, and every EC transition is accompanied by just one photon. Ans.: 1.15×10^9 yr.

3. Compute the number of grams and the number of radioactive atoms contained in 1 mc of (a) radiosodium (Na²⁴, T = 14.8 hr); (b) radiophosphorus (P²², T = 14.5 days); and (c) radium (Ra²²⁴, T = 1,620 yr). Ans.: mass,

1.1 × 10-10 g of Na24; 3.5 × 10-9 g of P22; 0.0010 g of Ra.

4. (a) A radioactive substance has a mean life τ sec, an activity of a_1 disintegrations per second at time t_1 , and an activity of a_2 at time t_2 . Show that the number of atoms $(A_1 - A_2)$ disintegrating between t_1 and t_2 is

$$A_1 - A_2 = \tau(a_1 - a_2)$$

(b) If the average energy per β ray of 12.6-hr iodine I¹³⁰ is 0.29 Mev, determine the β -ray energy in ergs liberated in 24 hr by an iodine source whose initial strength

(c) If this iodine is present in 2 g of thyroid tissue, determine the radiation is 1 mc. dose absorbed in 24 hr by the tissue, remembering that I rep (roentgen equivalent physical) corresponds to the absorption of 94 ergs per gram of tissue. Ans.: (b) 8.2×10^5 ergs; (c) 4,400 rep.

5. In 1 g of natural uranium,

(a) What is the activity of U^{238} , UX_1 , UX_2 , and U^{234} , in μ c? (b) What is the ratio of the activity of Uses to that of Uses?

(c) What is the number of spontaneous fissions per hour?

DATA: The decay series of U236 and U236 and half-periods are

$$\begin{array}{c}
a \\
22 U^{235} \xrightarrow{\alpha} & AcX^{223} \xrightarrow{\alpha} & AcX^{223} \xrightarrow{\alpha} & AcX^{223} \xrightarrow{\alpha} & AcX^{223} \xrightarrow{\alpha} & AcX^{223} \xrightarrow{\alpha} & AcX^{223} \xrightarrow{\alpha} & AcX^{223} & AcX^{2$$

In natural uranium, there is 1 atom of Uses per 139 atoms of Uses. The partial half-periods for spontaneous fission are

-periods for spontaneous hasion are

$$U^{231} = 8.0 \times 10^{15} \text{ yr}$$
 $U^{231} = 1.9 \times 10^{17} \text{ yr}$
 $U^{232} = 2 \times 10^{16} \text{ yr}$
 $U^{233} = 2 \times 10^{16} \text{ yr}$

Ans.: (a) 0.33 μc per gram U for each; (b) 0.046; (c) 25 fissions per hour per gram U (these are generally useful numbers, worth memorizing).

6. If an atom is known to exist at t = 0, what is its probability of decaying in the time interval Δt between t and $t + \Delta t$, if its decay constant is λ ? Under what restrictions does this general relationship reduce to simply $\lambda \Delta t$?

Ans.: $(1 - e^{-\lambda \Delta t})e^{-\lambda t}$; reduces when $\lambda t \ll 1$ and $\lambda \Delta t \ll 1$.

§2]

[CH. 15

2. Radioactive-s

In a number of which is also radio sented by

where λ₄ is the de constant of atoms the number of at The limiting case

a. The Genera any time t, the acti of change dB/dt, i supply of new ato: rate of loss of B the

If the only source t = 0, then

and, with these in

From this differen for B as a functio that the general s

In order to evalu dB/dt from Eq. (2

If this is to be va and therefore we

The coefficient ha special case in wh

Hence $h_B = -h_A$

[CH. 15

that of its product. After t_m the activity of the product must therefore exceed that of its parent.

d. Daughter Much Shorter-lived Than Parent. When the half-period of the daughter product is negligible compared with that of its parent, then Eq. (2.9) takes on a particularly simple form. Then $\lambda_4 \ll \lambda_B$, and Eq. (2.9) becomes

$$B\lambda_B = A\lambda_A(1 - e^{-\lambda_B t}) \tag{5.8}$$

The daughter activity $B\lambda_B$ increases according to the simple exponential growth curve governed by its own decay constant λ_B . This was the historically important case discovered by Rutherford and Soddy (R52) in the growth of ThX (T=3.64 days) from thorium (actually from RdTh, T=1.90 yr). Other important examples include the growth of radon in radium sources, etc. In these cases the equilibrium ratio of activities becomes substantially unity. Note then that

$$B\lambda_B = A\lambda_A \quad \text{for } t \gg T_B \tag{5.9}$$

only if $T_A \gg T_B$. This condition is spoken of classically as secular equilibrium.

6. Yield of a Radioactive Nuclide Produced by Nuclear Bombardment

Consider any nuclear reaction which results in the production of a radioactive nuclide, e.g.,

$$_{11}Na^{22} + H^2 \rightarrow H^1 + _{11}Na^{24}$$

In this reaction the number of target atoms of Na²² which are accessible to the deuteron beam can be called A_0 . The probability of transforming one of these atoms into Na²⁴ in unit time can be called λ_A . Then $A_0\lambda_A$ is the rate at which new atoms of Na²⁴ are produced. We see that the target is to be treated mathematically as though it were a parent source, having an activity $A_0\lambda_A$, and producing a radioactive substance B. Thus the scheme

$$A \xrightarrow{\lambda_a} B \xrightarrow{\lambda_B} C$$

represents the reactions

$$Na^{23} \xrightarrow{(d,p)} Na^{24} \xrightarrow{\beta^-} Mg^{24}$$

The probability λ_A of producing the (d,p) reaction is very small, but the number of target atoms A_0 is very large. Hence, mathematically,

$$A_0\lambda_A$$
 is finite $\lambda_A \rightarrow 0$ $A_0 \rightarrow \infty$

Usually, a negligible fraction of the atoms of the target is transformed so that the number of residual target atoms, $A = A_0 e^{-\lambda_A t}$, is effectively equal to A_0 . However, in some exceptional instances a measurable fraction of the target may be consumed, such as in the production of plutonium through intense and prolonged neutron irradiation of uranium.

§6]

In the Na²¹
after a uniform
for the growth of
lived parent.

The yield Y activity (not ato ment conditions yield of Na²⁴ f 14-Mev deutero

The yield is tant but specia and is equal to

For $B\lambda_B$ we can let us use instead in general, the

Thus in the great of the "parent'

Note that the y activity per unit then written as

The maximum athe maximum under the concidenterons,

 Y_{τ_B}

and this is the "bombardment, one-half this ul (6.1) shows that more than one decays almost production and periods $(e^{-\lambda t} =$